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REVIEW

Traditional Chinese medicine in COVID-19

Ming Lyu^{a,b,†}, Guanwei Fan^{c,†}, Guangxu Xiao^{a,†}, Taiyi Wang^d, Dong Xu^a, Jie Gao^e, Shaoqin Ge^e, Qingling Li^f, Yuling Ma^d, Han Zhang^a, Jigang Wang^b, Yuanlu Cui^{a,*}, Junhua Zhang^{a,*}, Yan Zhu^{a,*}, Boli Zhang^{a,*}

^aState Key Laboratory of Component-based Chinese Medicine, Tianjin University of Traditional Chinese Medicine, Tianjin 301617, China

^bArtemisinin Research Center, Institute of Chinese Materia Medica, China Academy of Chinese Medical Sciences, Beijing 100700, China

^cNational Clinical Research Center for Chinese Medicine Acupuncture and Moxibustion, First Teaching Hospital of Tianjin University of Traditional Chinese Medicine, Tianjin 300193, China

^dOxford Chinese Medicine Research Centre, University of Oxford, Oxford OX1 3PT, UK
^eCollege of Traditional Chinese Medicine, Hebei University, Baoding 071002, China
^fInstitute of Basic Medicine and Cancer, the Cancer Hospital of the University of Chinese

Academy of Sciences (Zhejiang Cancer Hospital), Chinese Academy of Sciences, Zhejiang 310022, China

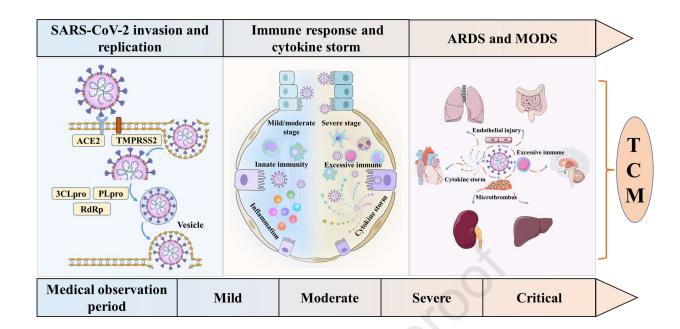
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*Corresponding authors.

E-mail addresses: yanzhu.harvard@icloud.com (Yan Zhu), zhangbolipr@163.com (Boli Zhang), zjhtcm@foxmail.com (Junhua Zhang), cuiyl@tju.edu.cn (Yuanlu Cui).

[†]These authors made equal contributions to this work.

Running title: TCM in COVID-19



Abstract COVID-19 pandemic caused by severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) has spread across the globe, posing an enormous threat to public health and safety. Traditional Chinese medicine (TCM), in combination with Western medicine (WM), has made important and lasting contributions in the battle against COVID-19. In this review, updated clinical effects and potential mechanisms of TCM, presented in newly recognized three distinct phases of the disease, are summarized and discussed. By integrating the available clinical and preclinical evidence, the efficacies and underlying mechanisms of TCM on COVID-19, including the highly recommended three Chinese patent medicines and three Chinese medicine formulas, are described in a panorama. We hope that this comprehensive review not only provides a reference for health care professionals and the public to recognize the significant contributions of TCM for COVID-19, but also serves as an evidence-based in-depth summary and analysis to facilitate understanding the true scientific value of TCM.

KEY WORDS COVID-19; SARS-CoV-2; Traditional Chinese medicine; Clinical evidence; Potential mechanism; Viral infection; Cytokine storm; Multiple organ dysfunction

1. Introduction

The outbreak and spread of coronavirus disease-19 (COVID-19) caused by severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) has inflicted immense losses on human lives and properties all over the world. Globally, as of August 7, 2021, there have been more than two hundred million confirmed COVID-19 cases, including more than four million of deaths (WHO, https://covid19.who.int/). SARS-CoV-2 is an enveloped, single-stranded, positive-sense, β -coronavirus RNA virus that belongs to the subfamily Coronavirinae, family Coronavirdiae, order Nidovirales. It shares about 79.6% identity of genome sequence with SARS-CoV and 96% similarity with bat coronavirus at the whole-genome level^{1,2}. SARS-CoV-2 is transmitted from

person to person *via* respiratory droplets, high concentration of aerosols, and occasionally feces or urine. Currently, no approved specific anti-viral drug is recommended to defeat COVID-19, which may lead to acute respiratory distress syndrome (ARDS), multiple organ dysfunction syndrome (MODS), and even death.

It is well documented that traditional Chinese medicine (TCM) has accumulated abundant clinical experience and effective prescriptions to control and treat infectious diseases in about 500 epidemics occurred in China over more than 3000 years in the past³. The combined therapy of TCM and Western medicine (WM) had significantly reduced mortality, shortened duration of fever, decreased chest radiograph abnormalities, and relieved secondary fungal infections among patients receiving glucocorticoids in combating severe acute respiratory syndrome (SARS)⁴. Owning to the positive role of TCM in treating previous coronavirus pneumonias such as SARS, middle east respiratory syndrome (MERS), and other epidemic diseases⁴⁻⁹, the National Health Commission of China recommended to use TCM as one of the strategies for COVID-19 remedy. This epidemic was deemed as the category of "pestilence" with the pathological characteristics of "dampness, heat, toxin, deficiency, and stasis" under TCM theory 10-12. Over the past year, TCM achieved remarkable efficacy in treating patients at all stages infected with SARS-CoV-2 in China. Typical clinical characteristics contain clinical manifestations, laboratory findings, and chest imaging features, as well as the pathogenesis of SARS-CoV-2 infection and therapeutic targets including SARS-CoV-2 invasion and replication, immune response, and cytokine storm, ARDS and MODS were outlined in published papers. In this review, the therapeutic efficacities and pharmacological mechanisms of TCM for this epidemic disease were systematically documented and discussed, aiming at displaying an in-depth understanding of TCM against COVID-19.

2. TCM in the treatment of COVID-19

2.1. Understanding COVID-19 in TCM theory

In the theory of TCM, COVID-19 is deemed as the category of "dampness-toxin pestilence" The distinct disease stages of TCM treatment can be divided into mild, moderate, severe, and critical. The main patterns in mild stage are cold-damp constraint and damp-heat accumulation in the lung, where dispersing lung and removing pathogenic factors, and resolve turbidity with aroma are needed; The main patterns in moderate stage are damp-toxin constraint in the lung and cold-damp obstructing the lung, where eliminating heat and dampness, detoxification, and invigorate spleen are needed; The main patterns in severe stage are epidemic toxin blocking the lung, blazing of both qi and yin, where tonifying qi and yin, ventilating lung qi, co-treatment of lung and intestines are needed. The main patterns in critical stage are internal blockage and external desertion, where tonifying qi and preventing exhaustion, cool blood and nourishing yin, and restore consciousness are needed $^{13-15}$. Syndrome differentiation is one of the most important principles for TCM to treat COVID-19.

2.2. The recommended TCMs for distinct stages of COVID-19 treatment

According to the officially issued 7th and 8th trial version of *Diagnosis and Treatment Protocol for COVID-19 in China* and other references^{14,16–23}, there are more than 18 recommended TCMs to prevent and treat COVID-19, covering from medical observation period (suspected cases) to clinical treatment period (confirmed cases) including distinct disease stages of mild, moderate, severe, and critical, as shown in Fig. 1. Among them, the highly recommended three Chinese patent medicines (CPMs) are Jinhua Qinggan granules, Lianhua Qingwen capsule (granules), and Xuebijing injection, and three Chinese medicine formulas are Qingfei Paidu decoction, Huashi Baidu formula, and Xuanfei Baidu formula, with proven efficacies in treating COVID-19^{24,25}. Jinhua Qinggan granules clear heat and detoxifying, and diffuse the lung. It is composed of 12 herbal medicines originating from Maxingshigan–Yinqiaosan formula, which could shorten time to fever resolution in patients with H1N1 influenza virus infection occurred in 2009²⁶. Lianhua Qingwen

capsule (granules), containing 13 herbal medicines and with a clinical indication for clearing heat, diffusing the lung, and detoxifying, was an innovative CPM for the treatment of SARS in 2003^{27,28}. Xuebijing injection, a five-herbal injection medicine and with a clinical indication for dissolving stasis and detoxifying, was derived from a modified Xuefu Zhuyu decoction and was developed and marketed during SARS. The Chinese medicine formula Qingfei Paidu decoction consists of 21 herbal medicines from five classic formulas of Treatise on Febrile Diseases. It clears the lung and calm panting, and is the first recommended universal treatment formula for all stages from mild to critical of COVID-19^{25,29}. Huashi Baidu formula is composed of 14 medicinal herbs. It serves to clearing heat and detoxifying, removing dampness, mainly suitable for the treatment of mild, moderate, and severe COVID-19 patients 30,31. Xuanfei Baidu formula is derived from classic formulas including Maxing Shigan decoction and Maxing Yigan decoction, and is composed of 13 medicinal herbs. It detoxifies and removes blood stasis, diffuses the lung, removes dampness, clears heat, and is mainly applicable to treat mild and moderate COVID-19 patients³². Beyond the above mentioned medicines and formulas, Chinese herbal injections, including Xiyanping injection, Reduning injection, Tanreging injection, Shenfu injection, Shengmai injection, and Shenmai injection, were more suitable as supplemental treatments for severe or critical COVID-19 cases with their advantages of fast absorption, high bioavailability, and clearer ingredients in contrast to orally administrated $TCMs^{33-35}$.

2.3. Clinical evidence of TCM for COVID-19

A total of 40 representative clinical trials, including 11 randomized controlled trials (RCTs), 16 retrospective cohort studies (RCSs), 5 multi-center clinical observations, and 8 others were completed and summarized^{27,28,30–32,36–70}. According to the available clinical data, integrated TCM and WM exhibited several clinical advantages in COVID-19 treatment, including the outcomes of 1) clinical manifestations, 2) lung features, and 3) laboratory findings as shown in Table 1^{27,28,30–32,36–70}. Furthermore,

based on Table 1, the clinical evidence of TCM for typical characteristics of COVID-19 were analyzed and summarized in Table 2^{27,28,30–32,36–70}.

For mild or moderate stages: 1) the most typical clinical symptoms of fever, cough, and fatigue were relieved by Jinhua Qinggan granules³⁷, Lianhua Qingwen granules³⁹, Shufeng Jiedu capsule^{51,52}, Toujie Quwen granules⁵⁷, Lianhua Qingke granules⁵⁶, Xuanfei Baidu decoction³², and Maxing Shigan decoction⁵⁹; Lianhua Qingwen granules³⁸ and Shufeng Jiedu capsule⁵¹ improved the symptoms of short of breath and chest tightness; Jinhua Qinggan granules relieved the symptom of psychological anxiety³⁷, and Shufeng Jiedu capsule⁵³ improved the symptom of diarrhea. 2) Jinhua Qinggan granules³⁶, Shufeng Jiedu capsule⁵¹, and Toujie Quwen granules⁵⁷ promoted pneumonia inflammatory absorption or improve lung CT imaging. 3) Jinhua Qinggan granules³⁶, Lianhua Qingwen granules³⁹, Shufeng Jiedu capsule⁵², Xuanfei Baidu decoction³², and Toujie Quwen granules³⁷, Shufeng Jiedu capsule⁵¹, Toujie Quwen granules⁵⁷, Xuanfei Baidu decoction³², and Maxing Shigan decoction⁵⁹ reduced the level of CRP. Shufeng Jiedu capsule decrease the level of IL-6⁵⁴.

For severe or critical stages: 1) Xuebijing injection²¹³ and Qingfei Paidu decoction⁴⁵ improved the conditions of patients and reduced multiple organ dysfunction. 2) Xuebijing injection²¹³, Qingfei Paidu decoction⁴⁵, and Huashi Baidu formula³⁰ improved chest CT imaging or promoted lung lesions absorption; Chansu injection⁶¹ ameliorated the respiratory function and shorten the respiratory support step-down time. 3) Both Xuebijing injection⁴³ and Chansu injection⁶¹ improved the oxygenation index of PaO₂/FiO₂; Xuebijing injection⁴³ and Qingfei Paidu decoction⁴⁵ decreased the level of CRP, and increased WBC or lymphocyte count; In addition, Xuebijing injection reduced the level of inflammatory mediators of TNF- α , IP-10, MIP-1 β , and RANTES⁴²; Qingfei Paidu decoction decreased biochemical parameters of CK and LDH, and the level of blood urea nitrogen⁴⁵; Maxing Shigan decoction

increased CD4⁺ T and CD8⁺ T count⁵⁹; Huashi Baidu formula³⁰ decreased CRP, ESR, serum ferritin, and myoglobin level; Yidu–toxicity blocking lung decoction reduced the levels of IL-6 and TNF- α^{62} .

For all stages: 1) Qingfei Paidu decoction⁴⁶ and Qingfei Dayuan granules⁶³ ameliorated extensive adverse symptoms such as fever, cough, fatigue, chest tightness, and headache; Xuanfei Huazhuo decoction relieved the symptoms of cough, fever, sputum, diarrhea, fatigue, and loss of appetite⁶⁵. 2) Qingfei Paidu decoction⁴⁵, Qingfei Dayuan granules⁶³, Xuanfei Huazhuo decoction⁶⁵, Keguan-1⁶⁶, Qingfei Touxie Fuzheng recipe⁶⁷, Ganlu Xiaodu decoction⁶⁸, and Matrine injection⁶⁹ improved lung inflammation or lesions absorption. 3) Qingfei Paidu decoction⁴⁶, Qingfei Dayuan granules⁶³, Xuanfei Huazhuo decoction⁶⁵, Ganlu Xiaodu decoction⁶⁸, "Fei Yan No. 1"⁶⁴, Matrine and sodium chloride injection⁶⁹, and Diammonium glycyrrhizinate⁷⁰ increased WBC or lymphocyte count; Qingfei Touxie Fuzheng recipe⁶⁷ and Diammonium glycyrrhizinate⁷⁰ decreased the level of CRP, IL-6, and ESR; Qingfei Paidu decoction^{46,48} and Xuanfei Huazhuo decoction⁶⁵ reduced the level of CRP and ESR, and the biochemical parameters of AST and ALT. What's more, Qingfei Paidu decoction decreased the level of a thrombotic marker D-dimer⁴⁶.

A plentiful of clinical studies and analyses proved that integrated Chinese and Western medicine therapy are much better than pure use of WM for COVID-19^{71–80}. A recent systematic review and meta-analysis of RCTs involving 2275 patients revealed that integration of TCM and WM group was more effective than WM treatment alone in the indicators of clinical cure rate, conversion rate from mild to critical, length of hospital stay, total score of clinical symptoms, symptoms of fever, cough and fatigue, TCM syndrome, negative conversion rate of viral nucleic acid, inflammatory biomarkers of CRP and lung CT without significant difference in adverse effects^{81,82}. Another similar meta-analysis of RCTs including 1259 COVID-19 patients showed consistent results that TCM with WM treatment could improve the

amounts of severe and critical conversion, length of hospital stay, time of antipyretic, and resolution rate of fever, fatigue, and tachypnea⁸³.

In summary of clinical evidence, TCM is beneficial for treating COVID-19 in 1) relieving the typical symptoms of fever, cough, fatigue, dry throat, sore throat, sputum production, shortness of breath, myalgia, and diarrhea; shorting the duration of positive viral nucleic acid, reducing the time to symptom recovery and the progression to severe disease, and protecting against multi-organ injury; 2) improving the lung features including lung inflammatory absorption, CT imaging, lung injury, lung function, and oxygenation index; 3) regulating laboratory index including inflammatory and immune response related the count of WBC, lymphocyte, CD4⁺ T and CD8⁺ T, and the level of CRP, IL-6, TNF-α, and ESR, single or multi-organ injury related the level of procalcitonin, CK, LDH, ALT, and AST, and thrombosis related D-dimer level. Taking full advantage of integration of TCM and WM is one of the important reasons for the rapid containment of this epidemic in China. Additional high-quality RCTs are needed to demonstrate the effectiveness and adverse events of TCM in the treatment of COVID-19.

3. Potential mechanisms of TCM for COVID-19

The intervention of TCM for COVID-19 is greatly inspired by the successful experience of treating SARS in 2002–2003^{4–9}. SARS-CoV-2 is genetically more similar with SARS-CoV (about 80%) than MERS-CoV (about 50%)^{1,2,84}. According to sequence alignment and homology modeling, the critical targets of spike, 3C-like protease (3CLpro), papain-like protease (PLpro), and RNA- dependent RNA polymerase (RdRp) protease share 76%, 96%, 83%, 96% sequence similarity between SARS-CoV and SARS-CoV-2, respectively^{85–87}. We collected and summarized TCMs and their ingredients to reveal the specific mechanisms of TCM for the three phases of distinct disease stages of COVID-19^{42,88–164}, seen in Table 3^{42,88–120} and Table 4^{109,115–117,121-164}

3.1. Potential mechanisms of TCM for SARS-CoV-2 invasion and replication

Although the direct evidence is still lacking, increasing reports suggested that TCM resource holds great promises for agents against SARS-CoV-2 invasion and replication. Numerous efforts had been made to identify the antiviral effects of CPMs and herbals, as shown in Table 3. Lianhua Qingwen capsule with a half maximal inhibitory concentration (IC₅₀) of 411.2 μg/mL⁸⁹, Liu Shen capsule¹⁰⁷ with an IC₅₀ of 0.6 μg/mL, and Shuanghuanglian preparation with an IC₅₀ of 0.93–1.2 μL/mL were confirmed to inhibit SARS-CoV-2 replication in Vero E6 cells. In addition, Pudilan Xiaoyan oral liquid not only inhibited SARS-CoV-2-stimulated Vero E6 cells in vitro, but also showed the potential efficacy on SARS-CoV-2-infected human angiotensin converting enzyme-2 (hACE2) transgenic mice in vivo 108. Six herbal extracts of Cibotium barometz (Gouji), Gentiana scabra (Longdan), Dioscorea batatas (Shanyao), Cassia tora (Juemingzi), and Taxillus chinensis (Sangjisheng) were evaluated for the anti-SARS-CoV activities by screening out from more than 200 extracts of Chinese medicinal herbs using a Vero E6 cell-based assay 120. Among them, Gouji and Shanyao could significantly inhibit 3CLpro protease activity of SARS-CoV with IC_{50} values of 39 and 44 $\mu g/mL^{120}$. Another screen of 312 Chinese medicinal herb extracts discovered three widely used Chinese medicinal herbs of the family Polygonaceae involving Rheum officinale (Yaoyong Dahuang), Polygonum multiflorum (Heshouwu), and Caulis polygoni multiflori (Shouwuteng) blocking the interaction of SARS-CoV Spike protein and angiotensin converting enzyme 2 (ACE2) which may protect the host from virus invasion with the IC₅₀ values ranged from 1 to 10 g/mL¹¹⁷. It was not difficult to find that although several TCMs like Liu Shen capsule and Dahuang showed a good performance in suppressing viral replication or activity, more studies are still necessary to be implemented to reveal more receivable anti-viral CPMs and herbal extracts especially the recommended CPMs in vitro and in vivo.

Noticeably, a considerable number of ingredients derived from TCMs were found to have anti-viral invasion and anti-viral replication activities by targeting

diverse molecules, as seen in Table 4. The interaction between spike protein and ACE2, primed by serine protease transmembrane protease serine 2 (TMPRSS2), is the key step for SARS-CoV-2 host invasion. Emodin from Yaoyong Dahuang was able to inhibit S protein and ACE2 interaction with an IC₅₀ of 200 µmol/L¹¹⁷, while hesperidin from Citrus aurantium (Suancheng) was predicted to target the binding between spike RBD and ACE2 with high affinity¹²⁴. Besides, geniposide from Gardenia jasminoides (Zhizi) was found through virtual screening of 2140 compounds with pharmacophoric features, which could target the active site residues of TMPRSS2 with a binding energy score of -14.69, and is even greater than that of the standard inhibitor of camostat mesylate 126. Seven isolated tanshinones derived from Salvia miltiorrhiza (Danshen) including tanshinone IIA, tanshinone IIB, methyl tanshinonate, crytotanshinone, tanshinone I, dihydrotanshinone I, and rosmariquinone showed marked inhibitory activities to both proteases of 3CLpro and PLpro 149. Particularly, dihydrotanshinone I exerted powerful effects with IC₅₀ values of 14.4 μmol/L regarding 3CLpro and 4.9 μmol/L regarding PLpro¹⁴⁹. Furthermore, crytotanshinone exhibited the most potent nanomolar level inhibitory activity toward PLpro with an IC₅₀ of 0.8 µmol/L¹⁴⁹. Baicalin and baicalein, the major bioactive ingredients of Shuanghuanglian preparation, were characterized as the first noncovalent and nonpeptidomimetic inhibitors of SARS-CoV-2 3CLpro, also possessed good anti-SARS-CoV-2 activity in Vero E6 cell-based system¹⁰⁹. What's more, celastrol^{143,144}, tingenone¹⁴³, xanthoangelol E¹⁵⁰, and hesperetin¹⁵¹ targeting 3CLpro, while hirsutenone¹⁵⁵, methyl tanshinoate, tanshinone I¹⁴⁹, xanthoangelol E¹⁵⁰, isobavachalcone, 4'-O-methylbavachalcone, psoralidin¹⁵⁷, and tomentin A-E¹⁵⁸ targeting PLpro, may have relatively strong anti-viral replication efficacy with IC₅₀ below or near 10 µmol/L. Notably, the well-known anti-malarial 165, anti-tumor 166, and immune modulation ¹⁶⁷ compound artemisinin from Artemisia apiacea (Oinghao), and its derivatives including arteannuin B, artesunate, dihydroartemisinin, arteether, and lumefantrine presented favorable anti-SARS-CoV-2 effects. Among these artemisinin

derivatives, arteannuin B showed the highest anti-viral potential with an IC₅₀ of 10.28 µmol/L, while lumefantrine exerted therapeutic promise owing to its high plasma and lung concentrations after multiple dosing. The deeper pharmacological mechanism analysis revealed that these two compounds acted at the post-entry step of SARS-CoV-2 infection¹³⁷. Significantly, lycorine from Lycoris radiata (Shisuan) had a powerful inhibitory effect on virus activity with an IC₅₀ of 15.7 nmol/L and may serve as a candidate for the development of new anti-SARS-CoV-2 drug in the treatment of COVID-19¹⁶⁴. In addition, a Vero E6 cell-based large-scale anti-SARS-CoV-2 activity of 1058 natural compounds were screened, and 17 newly discovered compounds showed strong anti-virus propagation effects with the IC₅₀ values ranging from 0.011 to 11.03 µmol/L. Among them, bufalin from toad venom (Chansu) exerted the antiviral effect with an IC₅₀ of 18 nmol/L by targeting the ion transport function of Na⁺/K⁺-ATPase¹³⁹. Theaflavin was predicted to exert anti-viral replication by inhibiting RdRp activity¹³⁰. The binding affinities with the critical proteins of a portion of ingredients presented above were also predicted by in silico screening and molecular docking 124,168. Whether these TCM ingredients could be used to combat COVID-19 need further in vitro and in vivo validation. Pharmacokinetic profiles including absorption, distribution, metabolism, and excretion (ADME) on the promising leads should be further studied.

3.2. Potential mechanisms of TCM for immune and inflammatory regulation

Antiviral monotherapy for patients hospitalized with COVID-19 is quite not enough, especially for severely and critically ill patients¹⁶⁹. Except for the broad-spectrum antiviral activity, TCM process advantages in regulating immune response, suppressing cytokine storm through multiple avenues^{170–172}. Beyond inhibiting virus replication, Lianhua Qingwen capsule⁸⁹ and Liu Shen capsule¹⁰⁷ reduced proinflammatory cytokines production such as TNF-α, IL-6, MCP-1, and IP-10 in SARS-CoV-2 infected Huh-7 cells. In addition, Lianhua Qingwen capsule was analyzed to repair lung injury by modulating inflammatory process and cytokine storm⁹⁰. Maxing

Shigan decoction is the basic prescription of "three medicines and three formulas" apart from Xuebijing injection, was revealed to regulate immunity and reduce cytokine storm, as well as protect alveolar-capillary barrier of lung and relieve pulmonary edema by utilizing integrated network pharmacological approaches ¹⁰¹. As same as Maxing Shigan decoction, Oingfei Paidu decoction showed multiple immune regulation, anti-inflammation, and lung injury-repair activities with its main ingredients of baicalin, glycyrrhizin, hesperidin, and hyperoside by targeting proteins including TNF-α, IL-6, IL-10, and CCL2⁹³⁻⁹⁶. Furthermore, several ingredients such as baicalin and glycyrrhizin of Qingfei Paidu decoction could inhibit platelet aggregation⁹⁶. Dayuanyin is the basic formula of Qingfei Dayuan granules that might process an anti-inflammatory and immunoregulatory effects via acting on IL-6, IL-1\beta, and MCP-1, with its ingredients containing kaempferol, isoflavone, and formononetin^{63,104}. Glycyrrhizin is an anti-viral agent and clinically used antiinflammatory ingredient from Glycyrrhiza uralensis (Gancao) was determined to elevate immunity and suppress inflammatory stress through T cell receptor and VEGF signaling pathways 141,159,173. Matrine was not only predicted to suppress host cell apoptosis and inflammation by targeting the TNF-α, IL-6, and CASP3 in the TNF signaling, but also validated to reduce lung tissue damage and lung index by decreasing the production of IL-6, IL-10, TNF- α , and IFN- γ , increasing the percentage of CD4⁺ T cells, CD8⁺ T cells, and B cells in peripheral blood, and lessening viral load in lung tissue in a mouse model combining human coronavirus pneumonia with cold–dampness pestilence attacking the lung 112,113. Although systems pharmacology is a convenient and effective tool to propose the mechanism of action of TCM at a holistic level, all the results above need to be further validated. IL-6 was considered as one of the most important molecules in cytokine storm 174-182. Administration with Dayuanvin reduced the level of IL-6 in mild, moderate, and even severe clinical stages of COVID-19¹⁰⁴. Besides, Shufeng Jiedu capsule⁵⁴, Yidutoxicity blocking lung decoction⁶², Qingfei Touxie Fuzheng recipe⁶⁷, and

diammonium glycyrrhizinate⁷⁰ were confirmed to decrease the level of IL-6 in COVID-19 patients, as seen in Table 1. Interestingly, except for the strong anti-SARS-CoV-2 activity¹³⁷, artemisinin and its derivatives regulated multiple immune cells including macrophage, monocyte, dendritic cell, and T cell to inhibit pro-inflammatory cytokine release and cytokine storm outbreak to protect tissues from injury¹⁸³ (Table 3).

3.3. Potential mechanisms of TCM for ARDS and MODS treatment

In contrast with WM therapy, TCM is adept at treating complications of COVID-19 such as ARDS and MODS which are likely caused by the concurrence of viral toxicity, endothelial damage, cytokine storm, excessive immune, and microthrombus holistically (Table 3). Xuebijing injection was certified to treat severe pneumonia, sepsis, coagulopathy, SIRS, and MODS, owing to its various effects on cytokine reduction, immunoregulation, microcirculation improvement, anti-coagulation, proangiogenesis, and neutralization of released bacterial cytotoxins^{42,184–189}. Xuebijing injection was able to improve the oxygenation index of PaO₂/FiO₂ and reduce the level of pro-inflammatory cytokines of TNF- α , IP-10, MIP-1 β , and RANTES in the treatment of COVID-19^{42,43}. It was also reported that Xuebijing injection could downregulate the expression of IL-6, IL-1, TLR4, MAPK, and NF-κB, maintain the balance of Tregs and Th17 cells in acute lung injury 190-193. Besides, Xuebijing injection processed the potential to alleviate liver damage, acute lung injury-induced left ventricular ischemia/reperfusion, sepsis-induced acute kidney injury, and sepsisinduced myocardial injury via inhibiting inflammation, apoptosis, and endothelial injury^{194–199}. Systems pharmacological analysis revealed that Qingfei Paidu decoction could protect multi-organ including nervous system, sensory system, digestive system, and circulatory system by regulating key enzymes, G protein-coupled receptors, ion channels, and transporters⁹⁶.

In the background of great demands for acute lung injury and ARDS therapy of COVID-19, more than one hundred of natural products from TCM with their potential

benefits and underlying mechanisms of anti-inflammation, antioxidant stress, antiapoptosis, and anti-pulmonary fibrosis were summarized and categorized. According to their chemical structures, these were divided into flavonoids (e.g., luteolin, baicalein), alkaloids (e.g., berberine, matrine), terpenoids (e.g., pogostone, andrographolide), polyphenols (e.g., honokiol, curcumin), quinonoids (e.g., emodin, shikonin), and other compounds (e.g., osthole, imperatorin)²⁰⁰. In addition, a systematic review and meta-analysis of 19 eligible RCTs including Tanreqing injection, Shengmai injection, Shenfu injection, Danshen injection, Reduning injection, and Xuebijing injection demonstrated that Chinese medicine injections were adjuvant therapy with great potential benefits for the treatment of ALI/ARDS³³. For example, based on the effects of inhibiting inflammatory cytokines of IL-6, IL-8, IL- 1β , and TNF- α , regulating immune, and elevating the oxygenation index of PaO₂, Tanreging injection was proved to improve lung injury, pulmonary infection, airway inflammation, and airway mucus hypersecretion²⁰¹⁻²⁰⁴. Reduning injection was demonstrated to prevent pulmonary neutrophil infiltration, lung injury and severe pneumonia which may attribute to downregulating IL-1 β , IL-18, TNF- α , NF- κ B, and pyrin domain containing 3 levels, lowering myeloperoxidase activities, and reducing reactive oxygen species production^{205–207}. Xiyanping injection, a famous Chinese medicinal preparation of andrographolide sulfonate, was reputed as one of the most effective alternatives to antibiotics, which has been widely used to ameliorate lung damage, bronchitis and community acquired pneumonia probably through inhibiting NF- κ B and MAPK-mediated inflammatory responses^{208,209}. Besides, Xiyanping injection and Reduning injection were used to treat diarrhea in children. Xiyanping injection could ameliorate colitis by inhibiting Th1/Th17 response in mice²¹⁰.

Cardiovascular disease is a high frequent comorbidity and complication of COVID-19. Three Chinese injection medicines including Shenfu injection, Shengmai injection, and Shenmai injection, have both pulmonary and cardiac protective effects. For instance, Shenfu injection is effective in the treatment of heart failure, myocardial

hypertrophy, cardiac arrest, myocardial ischemia-reperfusion injury, myocardial fibrosis, and acute viral myocarditis, partly through suppressing apoptosis and inflammation, improving microcirculation, reducing mitochondrial damage and coagulation-fibrinolysis disorders^{211–221}. Moreover, Shenfu injection has a protective effect on gastrointestinal tract and intestinal mucosa^{222,223}. Xingnaojing injection and Angong Niuhuang pill are different preparations share similar ingredients for stroke treatment in clinic. Both of them ameliorate cerebral ischemia/reperfusion injury, cerebral infarction, cerebral edema, blood–brain barrier disruption, and acute cerebral hemorrhage because of their benefits in brain microvascular endothelial cells, hippocampal and cortical neurons protection, and their anti-inflammation and anti-apoptosis effects^{224–231}.

3.4. Potential mechanisms of the representative and commonly used herbs in the treatment of COVID-19

Analyses of the main compositions of the "three medicines and three formulas" and other related literatures identified *Glycyrrhiza uralensis* (Gancao), *Ephedrae Herba* (Mahuang), *Semen Armeniacae Amarum* (Kuxingren), *Scutellaria baicalensis* (Huangqin), *Forsythiae Fructus* (Lianqiao), *Lonicera japonica* (Jingyinhua), *Rheum palmatum* (Dahuang), and *Artemisia annua* (Qinghao) as the representative and commonly used herbs for COVID-19^{3,81,232}. Herb—ingredient—target—function action network is established to elucidate the potential mechanisms of the frequently used herbs for COVID-19. In this relationship network, 8 commonly used herbs, 12 main ingredients, 10 key targets and 5 pivotal functions are involved, as shown in Fig. 2A. The portraits of commonly used herbs, chemical structures of ingredients, and main functions are illustrated in Fig. 2B.

Gut microbiome is involved in disease severity and host inflammatory and immune responses in COVID-19 patients²³³. It is worth noting that the anti-COVID-19 effects and mechanisms of TCM may be exerted *via* the gut-lung axis and mediated by gut microbiota^{234–236}. For example, short-term intervention of Qingfei

Paidu decoction dose-dependently regulates the host metabolism and gut microbiome in rats, indicating that altering gut microbiota composition may be part of the anti-COVID-19 mechanisms of Qingfei Paidu decoction²³⁷. It is of particularly significance to consider that the solubility and bioavailability of certain TCM ingredients, such as resveratrol, quercetin, baicalin, curcurmin, emodin, and tanshinone IIA, are limited, leading to poor absorption into the bloodstream after oral administration. These ingredients may exert their therapeutic effects though interplaying with gut microbiota²³⁸. For instance, resveratrol could also alleviate intestinal inflammation and oxidative damage by modulating the composition of gut microbiota in addition to the direct antiviral effect²³⁹. What's more, to improve the bioavailability, a nano-micellar form of curcumin was used to decrease IL-6 and IL- 1β expression and secretion in patients with COVID-19²⁴⁰.

In summary of preclinical evidence, the anti-COVID-19 effects and mechanisms of TCM include but not limited to 1) inhibiting SARS-CoV-2 invasion and replication by targeting the key proteins of spike, ACE2, TMPRSS2, 3CLpro, PLpro, RdRp, and spike–ACE2 interaction; 2) regulating immune and inflammatory response by targeting inflammatory cytokines such as IL-1, TNF-α, and IL-8, and chemokines like CCL5, CCL2, and IP-10, which are secreted by monocytes, macrophages, dendritic cells, CD4⁺ T cells, and CD8⁺ T cells; 3) protecting against ARDS and MODS by suppressing the crosstalk of viral toxicity, endothelial damage, cytokine storm, excessive immune, and microthrombus by targeting IL-6, CRP, D-dimer, and procalcitonin.

Finally, by integrating the clinical evidence and potential mechanisms of TCM for COVID-19, a panorama is drawn in Fig. 3, hoping that the effect and mechanism of TCM for COVID-19 could be viewed and understood within a single framework.

4. Conclusions and perspectives

Although a great quantity of review articles have been published on the topic of TCM in COVID-19^{13,14,19,23,25,35,71,87,168,171,241–277}, our work offers something unique. 1) To

our knowledge, this is the first review of TCM on COVID-19 that integrates evidence-based scientific findings from bedside to bench with the most comprehensive and updated literatures. 2) The pathogenesis and penitential mechanisms of TCM remedy in three phases corresponding to distinct stages for COVID-19 are first systematically described and presented within a single panorama by integrating available clinical and fundamental evidence.

A valuable lesson learned from China's COVID-19 battle is that perseverance in combination of TCM and WM is the right and sensible choice^{71,249}. Looking ahead, several critical issues need to be addressed as we prepare to face similar or even more serious global health threats in the future. Firstly, as the pandemic continues to evolve, the pathogenesis of COVID-19 is not fully elucidated. It is reasonable to postulate that the crosstalk of viral toxicity, endothelial damage, cytokine storm, excessive immune, and microthrombus are essential contributors for severely or critically ill patients with COVID-19, which need to be validated further. Secondly, due to a lack of in-depth understanding, there are still some skepticisms on the validity of treating COVID-19 with TCM^{278–280}. More RCTs with high accuracy, clinical safety, rigorous design, and large sample, as well as in-depth mechanistic explorations with compatibility principal should be conducted to provide more reliable evidence for TCM in COVID-19 intervention, especially for the highly recommended three CPMs and three Chinese medicine formulas. Thirdly, the rehabilitative effects of TCM ought to be continuous concerned and long-term medical observed for the COVID-19 patients in recovery phase, especially for the aged. A recent paper published in *The Lancet* on 6-month consequences of 1733 COVID-19 patients revealed that those with severe disease discharged from hospital showed common syndromes of fatigue or muscle weakness, sleep difficulties, and anxiety or depression^{281,282}. Meanwhile, a comparison of 425 non-treatment with 143 TCM-treated COVID-19 patients post discharge showed that TCM was beneficial for decreasing IL-6 and procalcitonin, and increasing red blood cell, hemoglobin, and platelet count²⁸³.

Overall, the purpose of this review is to scientifically and systematically evaluate the roles of TCM in combating COVID-19. The efficacies and potential mechanisms of TCM remedy in three phases of distinct stages of COVID-19 are discussed and presented comprehensively within a single panorama by integrating available clinical and preclinical evidence. Finally, although the availability of anti-COVID-19 vaccines and a global vaccination program have brought great hope for the ultimate control of the disease, threat of viral variants and new epidemics still exit. Therefore, it is of scientific value to historically and objectively summarize the contribution of TCM during the pandemic, which could be deployed in the future to combat against COVID-19 and other infectious diseases around the world.

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Author contributions

Boli Zhang, Yan Zhu, Junhua Zhang, Yuanlu Cui, and Jigang Wang conceived, designed, and revised the manuscript; Ming Lyu, Guanwei Fan, and Guangxu Xiao wrote and revised the manuscript; Taiyi Wang, Dong Xu, Jie Gao, Shaoqin Ge, Qinglin Li, Yuling Ma, and Han Zhang revised the manuscript and discussed interpretation.

Conflicts of interest

The authors declare no conflicts of interest.

References

- 1. Lu R, Zhao X, Li J, Niu P, Yang B, Wu H, et al. Genomic characterisation and epidemiology of 2019 novel coronavirus: implications for virus origins and receptor binding. *Lancet* 2020;**395**:565–74.
- 2. Zhou P, Yang X L, Wang X G, Hu B, Zhang L, Zhang W, et al. A pneumonia outbreak associated with a new coronavirus of probable bat origin. *Nature* 2020;579:270–3.
- 3. Luo H, Tang QL, Shang YX, Liang SB, Yang M, Robinson N, et al. Can Chinese medicine be used for prevention of corona virus disease 2019 (COVID-19)? A review of historical classics, research evidence and current prevention programs. *Chin J Integr Med* 2020;**26**:243–50.
- 4. Liu J, Manheimer E, Shi Y, Gluud C. Chinese herbal medicine for severe acute respiratory syndrome: a systematic review and meta-analysis. *J Altern Complement Med* 2004;**10**:1041–51.
- 5. Leung PC. The efficacy of Chinese medicine for SARS: a review of Chinese publications after the crisis. *Am J Chin Med* 2007;**35**:575–81.
- 6. International expert meeting on the treatment of SARS by traditional Chinese medicine, and the integration of traditional Chinese medicine with Western medicine. SARS: clinical trials on treatment using a combination of traditional Chinese medicine and Western medicine: report of the WHO international expert meeting to review and analyse clinical reports on combination treatment for SARS, 8–10 October 2003, Beijing, People's Republic of China. World Health Organization. Available from: https://apps.who.int/iris/handle/10665/43029.
- 7. Lau TF, Leung PC, Wong EL, Fong C, Cheng KF, Zhang SC, et al. Using herbal medicine as a means of prevention experience during the SARS crisis. *Am J Chin Med* 2005;**33**:345–56.

- 8. Hsu CH, Hwang KC, Chao CL, Chang SG, Ker CC, Chien LC, et al. The lesson of supplementary treatment with Chinese medicine on severe laboratory-confirmed SARS patients. *Am J Chin Med* 2006;**34**:927–35.
- 9. Chen Z, Nakamura T. Statistical evidence for the usefulness of Chinese medicine in the treatment of SARS. *Phytother Res* 2004;**18**:592–4.
- Zhao YS, Hou XY, Gao ZH, Wang T. Research on medication for severe type of COVID-19 based on Huashi Baidu prescription. *Chin Arch Trad Chin Med* 2020;38:14–7.
- 11. Zhao ZH, Zhou Y, Li WH, Huang QS, Tang ZH, Li H. Analysis of traditional Chinese medicine diagnosis and treatment strategies for COVID-19 based on "the diagnosis and treatment program for coronavirus disease-2019" from Chinese authority. *Am J Chin Med* 2020;**48**:1035–49.
- 12. Tong XL, Li XY, Zhao LH, Li QW, Yang YY, Lin YQ, et al. Discussion on traditional Chinese medicine prevention and treatment strategies of coronavirus disease 2019 (COVID-19) from the perspective of "cold–dampness pestilence". *J Ttadit Chin Med* 2020;**61**:465–70.
- 13. Leung ELH, Pan HD, Huang YF, Fan XX, Wang WY, He F, et al. The scientific foundation of Chinese herbal medicine against COVID-19. *Engineering* (Beijing) 2020;6:1099–107.
- 14. Luo H, Gao Y, Zou J, Zhang S, Chen H, Liu Q, et al. Reflections on treatment of COVID-19 with traditional Chinese medicine. *Chin Med* 2020;**15**:94.
- Zheng WK, Zhang JH, Yang FW, Huang M, Miao Q, Qi WS, et al. Treatment of coronavirus disease 2019 (COVID-19) from perspective of dampness–toxicity plagues. *J Tradit Chin Med* 2020;61:1024–28.
- 16. Chinese Association of Integrated Traditional and Western Medicine. Expert consensus on prevention and treatment of COVID-19 by integrating traditional Chinese and Western medicine. Chin J Integr Tradit West Med 2020;40:1413–23.
- 17. Li YX, Li J, Zhang Y, Tian YP, Zhang YG, Jin RJ, et al. Clinical practice

- guidelines and experts' consensuses for treatment of coronavirus disease 2019 (COVID-19) patients with Chinese herbal medicine: a systematic review. *Chin J Integr Med* 2020;**26**:786–93.
- 18. Lee BJ, Lee JA, Kim KI, Choi JY, Jung HJ. A consensus guideline of herbal medicine for coronavirus disease 2019. *Integr Med Res* 2020;**9**:100470.
- 19. Chan KW, Wong VT, Tang SCW. COVID-19: An update on the epidemiological, clinical, preventive and therapeutic evidence and guidelines of integrative Chinese–Western medicine for the management of 2019 novel coronavirus disease. *Am J Chin Med* 2020;**48**:737–62.
- 20. Liang N, Li H, Wang J, Jiao L, Ma Y, Wang X, et al. Development of rapid advice guidelines for the treatment of coronavirus disease 2019 with traditional Chinese medicine. *Am J Chin Med* 2020;**48**:1511–21.
- Ho LTF, Chan KKH, Chung VCH, Leung TH. Highlights of traditional Chinese medicine frontline expert advice in the China national guideline for COVID-19. Eur J Integr Med 2020;36:101116.
- 22. Liang N, Ma Y, Wang J, Li H, Wang X, Jiao L, et al. Traditional Chinese medicine guidelines for coronavirus disease 2019. *J Tradit Chin Med* 2020;**40**:891–6.
- 23. Qiu Q, Huang Y, Liu X, Huang F, Li X, Cui L, et al. Potential therapeutic effect of traditional Chinese medicine on coronavirus disease 2019: a review. *Front Pharmacol* 2020;**11**:570893.
- 24. Wang J, Qi F. Traditional Chinese medicine to treat COVID-19: the importance of evidence-based research. *Drug Discov Ther* 2020;**14**:149–50.
- 25. Li Q, Wang H, Li X, Zheng Y, Wei Y, Zhang P, et al. The role played by traditional Chinese medicine in preventing and treating COVID-19 in China. *Front Med* 2020;**14**:681–8.
- 26. Wang C, Cao B, Liu QQ, Zou ZQ, Liang ZA, Gu L, et al. Oseltamivir compared with the Chinese traditional therapy maxingshigan-yinqiaosan in the treatment of

- H1N1 influenza: a randomized trial. Ann Intern Med 2011;155:217–25.
- 27. Hu K, Guan WJ, Bi Y, Zhang W, Li L, Zhang B, et al. Efficacy and safety of Lianhuaqingwen capsules, a repurposed Chinese herb, in patients with coronavirus disease 2019: a multicenter, prospective, randomized controlled trial. *Phytomedicine* 2020;**16**:153242.
- 28. Xiao M, Tian J, Zhou Y, Xu X, Min X, Lv Y, et al. Efficacy of Huoxiang Zhengqi dropping pills and Lianhua Qingwen granules in treatment of COVID-19: a randomized controlled trial. *Pharmacol Res* 2020;**161**:105126.
- 29. Ren W, Ma Y, Wang R, Liang P, Sun Q, Pu Q, et al. Research advance on Qingfei Paidu decoction in prescription principle, mechanism analysis and clinical application. *Front Pharmacol* 2020;**11**:589714.
- 30. Huang L. Efficacy and safety assessment of severe COVID-19 patients with Chinese medicine: a retrospective case series study at early stage of the COVID-19 epidemic in Wuhan, China. *J Ethnopharmacol* 2021;**30**:113888.
- 31. Shi N, Guo L, Liu B, Bian Y, Chen R, Chen S, et al. Efficacy and safety of Chinese herbal medicine *versus* lopinavir–ritonavir in adult patients with coronavirus disease 2019: a non-randomized controlled trial. *Phytomedicine* 2020;**81**:153367.
- 32. Xiong WZ, Wang G, Du J, Ai W. Efficacy of herbal medicine (Xuanfei Baidu decoction) combined with conventional drug in treating COVID-19: a pilot randomized clinical trial. *Integr Med Res* 2020;**9**:100489.
- 33. Chen YB, Liu Q, Xie H, Yin SM, Wu L, Yu XH, et al. Is Chinese medicine injection applicable for treating acute lung injury and acute respiratory distress syndrome? A systematic review and meta-analysis of randomized controlled trials. *Chin J Integr Med* 2019;**6**:857–66.
- 34. Song P, Zhao L, Li X, Su J, Jiang Z, Song B, et al. Interpretation of the traditional Chinese medicine portion of the diagnosis and treatment protocol for corona virus disease 2019 (Trial Version 7). *J Tradit Chin Med* 2020;**40**:497–

508.

- 35. Zhang D, Zhang B, Lv JT, Sa RN, Zhang XM, Lin ZJ. The clinical benefits of Chinese patent medicines against COVID-19 based on current evidence. *Pharmacol Res* 2020;**157**:104882.
- 36. Liu Z, Li X, Gou C, Li L, Luo X, Zhang C, et al. Effect of Jinhua Qinggan granules on novel coronavirus pneumonia in patients. *J Tradit Chin Med* 2020;**40**:467–72.
- 37. Duan C, Xia WG, Zhen CJ, Sun GB, Li ZL, Li QL, et al. Clinical observation of Jinhua Qinggan granules in treating pneumonia infected by COVID-19. *J Tradit Chin Med* 2020;**61**:1473–7.
- 38. Lu RB, Wang WJ, Li X. Clinical observation on Lianhua Qingwen granules combined with Western medicine conventional therapy in the treatment of 63 suspected cases of COVID-19. *J Tradit Chin Med* 2020;**61**:655–9.
- 39. Yu P, Li YZ, Wan SB, Wang Y. Efficacy of Lianhua Qingwen granules combined with arbidol in the treatment of mild novel coronavirus pneumonia. *Chin J Chin Meter Med* 2020;55:1042–5.
- 40. Cheng ZD, Li Y. Clinical effectiveness and case analysis in 54 NCP patients treated with Lanhua Qingwen Granules. *World Chin Med* 2020;**15**:150–4.
- 41. Liu L, Shi F, Tu P, Chen C, Zhang M, Li X, et al. Arbidol combined with the Chinese medicine Lianhuaqingwen capsule *versus* arbidol alone in the treatment of COVID-19. *Medicine (Baltimore)* 2021;**100**:e24475.
- 42. Ma Q, Qiu M, Zhou H, Chen J, Yang X, Deng Z, et al. The study on the treatment of Xuebijing injection (XBJ) in adults with severe or critical Corona Virus Disease 2019 and the inhibitory effect of XBJ against SARS-CoV-2. *Pharmacol Res* 2020;**160**:105073.
- 43. Wen L, Zhou Z, Jiang D, Huang K. Effect of Xuebijing injection on inflammatory markers and disease outcome of coronavirus disease 2019. *Chin Crit Care Med* 2020;**32**:426–9.

- 44. Zhang CY, Zhang S, Wang W, Jiang XQ, Zhang CY, Zhang S, et al. Clinical observation of Xuebijing in the treatment of COVID-19. *Chin J Hosp Pharm* 2020;**40**:964–7.
- 45. Xin S, Cheng X, Zhu B, Liao X, Yang F, Song L, et al. Clinical retrospective study on the efficacy of Qingfei Paidu decoction combined with Western medicine for COVID-19 treatment. *Biomed Pharmacother* 2020;**129**:110500.
- 46. Wang RQ, Yang SJ, Xie CG, Shen QL, Li MQ, Lei X, et al. Clinical observation of Qingfei Paidu decoction in the treatment of COVID-19. *Pharmacol Clin Chin Mater Med* 2020;**36**:13–8.
- 47. Li KY, An W, Xia F, Chen M, Yang P, Liao YL, et al. Observation on clinical effect of modified Qingfei Paidu Decoction in treatment of COVID-19. *Chin Herb Med* 2020;**51**:2046–9.
- 48. Yu XY, Zhang S, Yan FF, Su DZ. Comparison of clinical efficacy of Qingfei Paidu decoction combined with western medicine in 43 cases and single western medicine in 46 cases in the treatment of COVID-19. *J Shandong Univ Health Sci* 2020;**58**:47–53.
- 49. Sun YN, Lv WL, Li H, Xiao Y, Yang W, Yang HJ, et al. A multicenter clinical study on the treatment of 295 COVID-19 cases with Qingfei Paidu decoction. *J Tradit Chin Med* 2020;**62**:599–603.
- 50. Shi N, Liu B, Liang N, Ma Y, Ge Y, Yi H, et al. Association between early treatment with Qingfei Paidu decoction and favorable clinical outcomes in patients with COVID-19: a retrospective multicenter cohort study. *Pharmacol Res* 2020;**161**:105290.
- 51. Chen L, Liu F, Wu JH, Song HY, Xia JS, Sheng B, et al. Clinical efficacy of Shufeng Jiedu capsule combined with western medicine in treatment of common COVID-19 patients by retrospective analysis. *Chin J Exp Tradit Med Form* 2020;**26**:14–20.
- 52. Xiao Q, Jiang YJ, Wu SS, Wang Y, An J, Xu WP, et al. Analysis of the value of

- Shufeng Jiedu capsule combined with Arbidol in the treatment of mild new coronavirus pneumonia. *J Emerg Tradit Chin Med* 2020;**29**:756–8.
- 53. Qu XK, Hao SL, Ma JH, Wei GY, Song KY, Tang C, et al. Observation on clinical effect of Shufeng Jiedu capsule combined with arbidol hydrochloride capsule in treatment of COVID-19. *Chin Herb Med* 2020;**51**:1167–70.
- 54. Chen J, Lin S, Niu C, Xiao Q. Clinical evaluation of Shufeng Jiedu Capsules combined with umifenovir (arbidol) in the treatment of common-type COVID-19: a retrospective study. *Expert Rev Respir Med* 2020;**15**:257–65.
- 55. Tian J, Yan S, Wang H, Zhang Y, Zheng Y, Wu H, et al. Hanshiyi Formula, a medicine for SARS-CoV-2 infection in China, reduced the proportion of mild and moderate COVID-19 patients turning to severe status: a cohort study. *Pharmacol Res* 2020;**161**:105127.
- 56. Sun HM, Xu F, Zhang L, Wei C, Chen JY, Wang QX, et al. Study on the clinical efficacy of Lianhua Qingke granules in the treatment of mild and common COVID-19. *Chin J Exp Tradit Med Form* 2020;**26**:29–34.
- 57. Fu XX, Lin LP, Tan XH. Clinical observation on effect of Toujie Quwen granules in treatment of COVID-19. *Chin J Exp Tradit Med Form* 2020;**26**:44–8.
- 58. Yang MB, Dang SS, Huang S, Li YJ, Guo YL. Multi-center clinical observation of Reyanning mixture in treatment of COVID-19. *Chin J Exp Tradit Med Form* 2020;**26**:7–12.
- 59. Qu YF, Fang W, Jin YZ, Qin C, Niu XC, Zhang N, et al. Forty cases of common COVID-19 treated with modified ephedra and apricot kernel and gypsum and licorice decoction combined with Western medicine routine treatment. *Henan Tradit Chin Med* 2020;**40**:666–9.
- 60. Hu F, Guo AH, Huang L, Yu WX, Liu GF, Gao XS, et al. Multi-center clinical observation of honeysuckle oral liquid combined with Western medicine in the treatment of common COVID-19. *J Tradit Chin Med* 2020;**62**:510–5.
- 61. Hu F, Chen J, Chen H, Zhu J, Wang C, Ni H, et al. Chansu injection improves the

- respiratory function of severe COVID-19 patients. *medRxiv* 2020. Available from: https://doi.org/10.1101/2020.05.20.20107607.
- 62. Zhao J, Yang X, Wang C, Song S, Cao K, Wei T, et al. Yidu–toxicity blocking lung decoction ameliorates inflammation in severe pneumonia of SARS-COV-2 patients with Yidu–toxicity blocking lung syndrome by eliminating IL-6 and TNF-a. *Biomed Pharmacother* 2020;**129**:110436.
- 63. Ba YM, Wang LQ, Li WN, Li M, Tao R, Zuo XH, et al. Multi center clinical study on 451 cases of COVID-19 treated with 'Pneumonia No.1 Formula'. *World Chin Med* 2020;**15**:1962–6.
- 64. Ai Z, Zhou S, Li W, Wang M, Wang L, Hu G, et al. "Fei Yan No. 1" as a combined treatment for COVID-19: An efficacy and potential mechanistic study. *Front Pharmacol* 2020;**11**:581277.
- 65. Shi TF, Zhou GC, Zhang LY, Niu F, Ke YC, Zhou T, et al. Clinical efficacy of Xuanfei Huazhuo decoction on 40 cases of COVID-19. *Chin J Exp Tradit Med Form* 2020;**26**:26–31.
- 66. Wang JB, Wang ZX, Jing J, Zhao P, Dong JH, Zhou YF, et al. Exploring an integrative therapy for treating COVID-19: a randomized controlled trial. *Chin J Integr Med* 2020;**26**:648–55.
- 67. Ding XJ, Zhang Y, He DC, Zhang MY, Tan YJ, Yu AR, et al. Clinical effect and mechanism of Qingfei Touxie Fuzheng recipe in the treatment of COVID-19. Herald Med 2020;39:640–4.
- 68. Chen L, Cheng ZQ, Liu F, Xia Y, Chen YG. Analysis of 131 cases of COVID-19 treated with Ganlu Xiaodu Decoction. *China J Chin Mater Med* 2020;**45**:2232–8.
- 69. Yang MW, Chen F, Zhu DJ, Li JZ, Zhu JL, Zeng W, et al. Clinical efficacy of matrine and sodium chloride injection in treatment of 40 cases of COVID-19. *China J Chin Mater Med* 2020;45:2221–31.
- 70. Xi JX, Xiang SL, Zhang HJ, Lan BQ, Chen XY. Clinical observation of Arbidol combined with diammonium lycyrrhizinate in the treatment of COVID-19. *Chin*

- J Hosp Phar 2020;**40**:1287–90.
- 71. Ni L, Chen L, Huang X, Han C, Xu J, Zhang H, et al. Combating COVID-19 with integrated traditional Chinese and Western medicine in China. *Acta Pharm Sin B* 2020;**10**:1149–62.
- 72. Xia WG, An CQ, Zhen CJ, Zhang JX, Huang M, Wang Y, et al. Clinical observation on 34 patients with novel coronavirus pneumonia (COVID-19) treated with integrated traditional Chinese and Western medicine. *J Tradit Chin Med* 2020;**61**:375–82.
- 73. Yang Q, Sun Q G, Jiang B, Xu HJ, Luo M, Xie P, et al. Retrospective clinical study on treatment of COVID-19 patients with integrated traditional Chinese and western medicine. *Chin Herb Med* 2020;**51**:2050–4.
- 74. Shu Z, Zhou Y, Chang K, Liu J, Min X, Zhang Q, et al. Clinical features and the traditional Chinese medicine therapeutic characteristics of 293 COVID-19 inpatient cases. *Front Med* 2020;14:760–75.
- 75. Zhang HT, Huang MX, Liu X, Zheng XC, Li XH, Chen GQ, et al. Evaluation of the adjuvant efficacy of natural herbal medicine on COVID-19: a retrospective matched case-control study. *Am J Chin Med* 2020;**48**:779–92.
- 76. Liu M, Gao Y, Yuan Y, Yang K, Shi S, Zhang J, et al. Efficacy and safety of integrated traditional Chinese and Western medicine for corona virus disease 2019 (COVID-19): a systematic review and meta-analysis. *Pharmacol Res* 2020;158:104896.
- 77. Ye YA. Guideline-based Chinese herbal medicine treatment plus standard care for severe coronavirus disease 2019 (G-CHAMPS): evidence from China. *Front Med (Lausanne)* 2020;7:256.
- 78. Zhang K, Tian M, Zeng Y, Wang L, Luo S, Xia W, et al. The combined therapy of a traditional Chinese medicine formula and Western medicine for a critically ill case infected with COVID-19. *Complement Ther Med* 2020;**52**:102473.
- 79. Chen G, Su W, Yang J, Luo D, Xia P, Jia W, et al. Chinese herbal medicine

- reduces mortality in patients with severe and critical coronavirus disease 2019: a retrospective cohort study. *Front Med* 2020;**14**:752–9.
- 80. Ang L, Song E, Lee HW, Lee MS. Herbal medicine for the treatment of coronavirus disease 2019 (COVID-19): a systematic review and meta-analysis of randomized controlled trials. *J Clin Med* 2020;**9**:1583.
- 81. Xiong X, Wang P, Su K, Cho WC, Xing Y. Chinese herbal medicine for coronavirus disease 2019: a systematic review and meta-analysis. *Pharmacol Res* 2020;**160**:105056.
- 82. Yu M, Zhang R, Ni P, Duan G. Chinese herbal medicine supplementation therapy on COVID-19. *Pharmacol Res* 2020;**160**:105181.
- 83. Pang W, Liu Z, Li N, Li Y, Yang F, Pang B, et al. Chinese medical drugs for coronavirus disease 2019: a systematic review and meta-analysis. *Integr Med Res* 2020;**9**:100477.
- 84. Chan JF, Kok KH, Zhu Z, Chu H, To KK, Yuan S, et al. Genomic characterization of the 2019 novel human–pathogenic coronavirus isolated from a patient with atypical pneumonia after visiting Wuhan. *Emerg Microbes Infect* 2020;9:221–36.
- 285. Morse JS, Lalonde T, Xu S, Liu WR. Learning from the past: possible urgent prevention and treatment options for severe acute respiratory infections caused by 2019-nCoV. *Chembiochem* 2020;**21**:730–8.
- 86. Liu C, Zhou Q, Li Y, Garner L V, Watkins SP, Carter LJ, et al. Research and development on therapeutic agents and vaccines for COVID-19 and related human coronavirus diseases. *ACS Cent Sci* 2020;**6**:315–31.
- 87. Li H, Yang L, Liu FF, Ma XN, He PL, Tang W, et al. Overview of therapeutic drug research for COVID-19 in China. *Acta Pharmacol Sin* 2020;**41**:1133–40.
- 88. Gong PY, Guo YJ, Li XP, Wang N, Gu J. Exploring active compounds of Jinhua Qinggan Granules for prevention of COVID-19 based on network pharmacology and molecular docking. *Chin Tradit Herb Drugs* 2020;**51**:1685–93.

- 89. Li RF, Hou YL, Huang JC, Pan WQ, Ma QH, Shi YX, et al. Lianhuaqingwen exerts anti-viral and anti-inflammatory activity against novel coronavirus (SARS-CoV-2). *Pharmacol Res* 2020;**156**:104761.
- 90. Zheng S, Baak JP, Li S, Xiao W, Ren H, Yang H, et al. Network pharmacology analysis of the therapeutic mechanisms of the traditional Chinese herbal formula Lian Hua Qing Wen in Corona virus disease 2019 (COVID-19), gives fundamental support to the clinical use of LHQW. *Phytomedicine* 2020;**79**:153336.
- 91. Xing Y, Hua YR, Shang J, Ge WH, Liao J. Traditional Chinese medicine network pharmacology study on exploring the mechanism of Xuebijing Injection in the treatment of coronavirus disease 2019. *Chin J Nat Med* 2020;**18**:941–51.
- 92. Zheng WJ, Yan Q, Ni YS, Zhan SF, Yang LL, Zhuang HF, et al. Examining the effector mechanisms of Xuebijing injection on COVID-19 based on network pharmacology. *BioData Min* 2020;**13**:17.
- 93. Chen J, Wang YK, Gao Y, Hu LS, Yang JW, Wang JR, et al. Protection against COVID-19 injury by Qingfei Paidu decoction *via* anti-viral, anti-inflammatory activity and metabolic programming. *Biomed Pharmacother* 2020;**129**:110281.
- 94. Yang R, Liu H, Bai C, Wang Y, Zhang X, Guo R, et al. Chemical composition and pharmacological mechanism of Qingfei Paidu Decoction and Ma Xing Shi Gan Decoction against coronavirus disease 2019 (COVID-19): *in silico* and experimental study. *Pharmacol Res* 2020;**157**:104820.
- 95. Zhang DH, Zhang X, Peng B, Deng SQ, Wang YF, Yang L, et al. Network pharmacology suggests biochemical rationale for treating COVID-19 symptoms with a traditional Chinese medicine. *Commun Biol* 2020;**3**:466.
- 96. Zhao J, Tian S, Lu D, Yang J, Zeng H, Zhang F, et al. Systems pharmacological study illustrates the immune regulation, anti-infection, anti-inflammation, and multi-organ protection mechanism of Qing-Fei-Pai-Du decoction in the treatment of COVID-19. *Phytomedicine* 2020;**85**:153315.

- 97. Tao Q, Du J, Li X, Zeng J, Tan B, Xu J, et al. Network pharmacology and molecular docking analysis on molecular targets and mechanisms of Huashi Baidu formula in the treatment of COVID-19. *Drug Dev Ind Pharm* 2020;**46**:1345–53.
- 98. Wang Y, Li X, Zhang JH, Xue R, Qian JY, Zhang XH, et al. Study on the mechanism of Xuanfei Baidu decoction for COVID-19 based on network pharmacology. *Chin J Chin Mater Med* 2020;**45**:2249–56.
- 99. Chen X, Yin YH, Zhang MY, Liu JY, Li R, Qu YQ. Investigating the mechanism of ShuFeng JieDu capsule for the treatment of novel coronavirus pneumonia (COVID-19) based on network pharmacology. *Int J Med Sci* 2020;**17**:2511–30.
- 100. Tao Z, Zhang L, Friedemann T, Yang G, Li J, Wen Y, et al. Systematic analyses on the potential immune and anti-inflammatory mechanisms of Shufeng Jiedu capsule against severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2)-caused pneumonia. *J Funct Foods* 2020;75:104243.
- 101. Wang YX, Ma JR, Wang SQ, Zeng YQ, Zhou CY, Ru YH, et al. Utilizing integrating network pharmacological approaches to investigate the potential mechanism of Ma Xing Shi Gan Decoction in treating COVID-19. Eur Rev Med Pharmacol Sci 2020;24:3360–84.
- 102. Li Y, Chu F, Li P, Johnson N, Li T, Wang Y, et al. Potential effect of Maxing Shigan decoction against coronavirus disease 2019 (COVID-19) revealed by network pharmacology and experimental verification. *J Ethnopharmacol* 2021;**271**:113854.
- 103. Han L, Wei XX, Zheng YJ, Zhang LL, Wang XM, Yang HY, et al. Potential mechanism prediction of Cold–Damp Plague Formula against COVID-19 via network pharmacology analysis and molecular docking. Chin Med 2020;15:78.
- 104. Ruan X, Du P, Zhao K, Huang J, Xia H, Dai D, et al. Mechanism of Dayuanyin in the treatment of coronavirus disease 2019 based on network pharmacology and molecular docking. *Chin Med* 2020;**15**:62.

- 105. Zhang XR, Li TN, Ren YY, Zeng YJ, Lv HY, Wang J, et al. The important role of volatile components from a traditional Chinese medicine Dayuan–Yin against the COVID-19 pandemic. *Front Pharmacol* 2020;11:583651.
- 106. Jia S, Luo H, Liu X, Fan X, Huang Z, Lu S, et al. Dissecting the novel mechanism of Reduning injection in treating Coronavirus Disease 2019 (COVID-19) based on network pharmacology and experimental verification. *J Ethnopharmacol* 2021;**273**:113871.
- 107. Ma Q, Pan W, Li R, Liu B, Li C, Xie Y, et al. Liu Shen capsule shows antiviral and anti-inflammatory abilities against novel coronavirus SARS-CoV-2 *via* suppression of NF-kappaB signaling pathway. *Pharmacol Res* 2020;**158**:104850.
- 108. Deng W, Xu Y, Kong Q, Xue J, Yu P, Liu J, et al. Therapeutic efficacy of Pudilan Xiaoyan Oral Liquid (PDL) for COVID-19 *in vitro* and *in vivo*. *Signal Transduct Target Ther* 2020;**5**:66.
- 109. Su HX, Yao S, Zhao WF, Li MJ, Liu J, Shang WJ, et al. Anti-SARS-CoV-2 activities *in vitro* of Shuanghuanglian preparations and bioactive ingredients. *Acta Pharmacol Sin* 2020;**41**:1167–77.
- 110. Lin H, Wang X, Liu M, Huang M, Shen Z, Feng J, et al. Exploring the treatment of COVID-19 with Yinqiao powder based on network pharmacology. *Phytother Res* 2021. Available from: https://doi.org/10.1002/ptr.7012.
- 111. Kong Q, Wu Y, Gu Y, Lv Q, Qi F, Gong S, et al. Analysis of the molecular mechanism of Pudilan (PDL) treatment for COVID-19 by network pharmacology tools. *Biomed Pharmacother* 2020;**128**:110316.
- 112. Peng W, Xu Y, Han D, Feng F, Wang Z, Gu C, et al. Potential mechanism underlying the effect of matrine on COVID-19 patients revealed through network pharmacological approaches and molecular docking analysis. *Arch Physiol Biochem* 2020. Available from: https://www.tandfonline.com/doi/full/10.1080/13813455.2020.1817944.
- 113. Sun J, Zhao RH, Guo SS, Shi YJ, Bao L, Geng ZH, et al. Effect of matrine

- sodium chloride injection on a mouse model combining disease with syndrome of human coronavirus pneumonia with cold-dampness pestilence attacking the lung. *Acta Pharmaceutica Sinica* 2020;**55**:366–73.
- 114. Li X, Lin H, Wang Q, Cui L, Luo H, Luo L. Chemical composition and pharmacological mechanism of shenfu decoction in the treatment of novel coronavirus pneumonia (COVID-19). *Drug Dev Ind Pharm* 2020;**46**:1947–59.
- 115. Sa-ngiamsuntorn K, Suksatu A, Pewkliang Y, Thongsri P, Kanjanasirirat P, Manopwisedjaroen S, et al. Anti-SARS-CoV-2 activity of *Andrographis paniculata* extract and its major component andrographolide in human lung epithelial cells and cytotoxicity evaluation in major organ cell representatives. *J Nat Prod* 2021;84:1261–70.
- 116. Liu H, Ye F, Sun Q, Liang H, Li C, Li S, et al. Scutellaria baicalensis extract and baicalein inhibit replication of SARS-CoV-2 and its 3C-like protease in vitro. J Enzym Inhib Med Ch 2021;36:497–503.
- 117. Ho TY, Wu SL, Chen JC, Li CC, Hsiang CY. Emodin blocks the SARS coronavirus spike protein and angiotensin-converting enzyme 2 interaction.

 *Antiviral Res 2007;74:92–101.
- 118. Lau KM, Lee KM, Koon CM, Cheung CS, Lau CP, Ho HM, et al. Immunomodulatory and anti-SARS activities of *Houttuynia cordata*. *J Ethnopharmacol* 2008;**118**:79–85.
- 119. Luo W, Su X, Gong S, Qin Y, Liu W, Li J, et al. Anti-SARS coronavirus 3C-like protease effects of *Rheum palmatum* L. extracts. *Biosci Trends* 2009;**3**:124–6.
- 120. Wen CC, Shyur LF, Jan JT, Liang PH, Kuo CJ, Arulselvan P, et al. Traditional Chinese medicine herbal extracts of *Cibotium barometz*, *Gentiana scabra*, *Dioscorea batatas*, *Cassia tora*, and *Taxillus chinensis* inhibit SARS-CoV replication. *J Tradit Complement Med* 2011;1:41–50.
- 121. Chen X, Wu Y, Chen C, Gu Y, Zhu C, Wang S, et al. Identifying potential anti-COVID-19 pharmacological components of traditional Chinese medicine

- Lianhuaqingwen capsule based on human exposure and ACE2 biochromatography screening. *Acta Pharm Sin B* 2021;**11**:222–36.
- 122. Liu X, Raghuvanshi R, Ceylan FD, Bolling BW. Quercetin and its metabolites inhibit recombinant human angiotensin-converting enzyme 2 (ACE2) activity. *J Agric Food Chem* 2020;**68**:13982–9.
- 123. Lv Y, Wang S, Liang P, Wang Y, Zhang X, Jia Q, et al. Screening and evaluation of anti-SARS-CoV-2 components from Ephedra sinica by ACE2/CMC-HPLC-IT-TOF-MS approach. *Anal Bioanal Chem* 2021;**413**:2995–3004.
- 124. Wu C, Liu Y, Yang Y, Zhang P, Zhong W, Wang Y, et al. Analysis of therapeutic targets for SARS-CoV-2 and discovery of potential drugs by computational methods. *Acta Pharm Sin B* 2020;**10**:766–88.
- 125. Haggag YA, El-Ashmawy NE, Okasha KM. Is hesperidin essential for prophylaxis and treatment of COVID-19 Infection?. *Med Hypotheses* 2020;**144**:109957.
- 126. Rahman N, Basharat Z, Yousuf M, Castaldo G, Rastrelli L, Khan H. Virtual screening of natural products against type II transmembrane serine protease (TMPRSS2), the priming agent of coronavirus 2 (SARS-CoV-2). *Molecules* 2020;**25**:2271.
- 127. Jin Z, Du X, Xu Y, Deng Y, Liu M, Zhao Y, et al. Structure of M(pro) from SARS-CoV-2 and discovery of its inhibitors. *Nature* 2020;**582**:289–93.
- 128. Jang M, Park YI, Cha YE, Park R, Namkoong S, Lee JI, et al. Tea polyphenols EGCG and theaflavin inhibit the activity of SARS-CoV-2 3CL-protease *in vitro*. *Evid Based Complement Alternat Med* 2020;**2020**:5630838.
- 129. Raj V, Park JG, Cho KH, Choi P, Kim T, Ham J, et al. Assessment of antiviral potencies of cannabinoids against SARS-CoV-2 using computational and *in vitro* approaches. *Int J Biol Macromol* 2021;**168**:474–85.
- 130. Lung J, Lin YS, Yang YH, Chou YL, Shu LH, Cheng YC, et al. The potential chemical structure of anti-SARS-CoV-2 RNA-dependent RNA polymerase. *J*

- *Med Virol* 2020;**92**:693–7.
- 131. Mhatre S, Srivastava T, Naik S, Patravale V. Antiviral activity of green tea and black tea polyphenols in prophylaxis and treatment of COVID-19: a review. *Phytomedicine* 2020;**85**:153286.
- 132. Jeon S, Ko M, Lee J, Choi I, Byun SY, Park S, et al. Identification of antiviral drug candidates against SARS-CoV-2 from FDA-approved drugs. *Antimicrob Agents Chemother* 2020;**64**:e00819-20.
- 133. Sand LVD, Bormann M, Alt M, Schipper L, Heilingloh CS, Todt D, et al. Glycyrrhizin effectively neutralizes SARS-CoV-2 *in vitro* by inhibiting the viral main protease. *Viruses* 2021;**13**:609.
- 134. Ellen ter BM, Dinesh N, Bouma EM, Troost B, Pol van de DPI, Ende Van der-Metselaar HH, et al. Resveratrol and pterostilbene potently inhibit SARS-CoV-2 infection *in vitro*. *Viruses* 2021;**13**:1335.
- 135. Ma Q, Li R, Pan W, Huang W, Liu B, Xie Y, et al. Phillyrin (KD-1) exerts antiviral and anti-inflammatory activities against novel coronavirus (SARS-CoV-2) and human coronavirus 229E (HCoV-229E) by suppressing the nuclear factor kappa B (NF-kappaB) signaling pathway. *Phytomedicine* 2020;**78**:153296.
- 136. Mishra CB, Pandey P, Sharma RD, Malik MZ, Mongre RK, Lynn AM, et al. Identifying the natural polyphenol catechin as a multi-targeted agent against SARS-CoV-2 for the plausible therapy of COVID-19: an integrated computational approach. *Brief Bioinform* 2021;22:1346–60.
- 137. Cao R, Hu H, Li Y, Wang X, Xu M, Liu J, et al. Anti-SARS-CoV-2 potential of artemisinins *in vitro*. *ACS Infect Dis* 2020;**6**:2524–31.
- 138. Fan HH, Wang LQ, Liu WL, An XP, Liu ZD, He XQ, et al. Repurposing of clinically approved drugs for treatment of coronavirus disease 2019 in a 2019-novel coronavirus-related coronavirus model. *Chin Med J (Engl)* 2020;**133**:1051–6.
- 139. Zhang ZR, Zhang YN, Li XD, Zhang HQ, Xiao SQ, Deng F, et al. A cell-based

- large-scale screening of natural compounds for inhibitors of SARS-CoV-2. Signal Transduct Target Ther 2020;5:218.
- 140. Clementi N, Scagnolari C, D'Amore A, Palombi F, Criscuolo E, Frasca F, et al. Naringenin is a powerful inhibitor of SARS-CoV-2 infection *in vitro*. *Pharmacol Res* 2021;**163**:105255.
- 141. Li R, Wu K, Li Y, Liang X, Lai KP, Chen J. Integrative pharmacological mechanism of vitamin C combined with glycyrrhizic acid against COVID-19: findings of bioinformatics analyses. *Brief Bioinform* 2021;22:1161–74.
- 142. Wang WX, Zhang YR, Luo SY, Zhang YS, Zhang Y, Tang C. Chlorogenic acid, a natural product as potential inhibitor of COVID-19: virtual screening experiment based on network pharmacology and molecular docking. *Nat Prod Res* 2021. Available from: https://www.tandfonline.com/doi/full/10.1080/14786419.2021.1904923.
- 2143. Ryu YB, Park SJ, Kim YM, Lee JY, Seo WD, Chang J S, et al. SARS-CoV 3CLpro inhibitory effects of quinone-methide triterpenes from *Tripterygium regelii*. *Bioorg Med Chem Lett* 2010;**20**:1873–6.
- 144. Habtemariam S, Nabavi SF, Berindan-Neagoe I, Cismaru CA, Izadi M, Sureda A, et al. Should we try the antiinflammatory natural product, celastrol, for COVID-19?. *Phytother Res* 2020;**34**:1189–90.
- 145. Wen CC, Kuo YH, Jan JT, Liang PH, Wang SY, Liu HG, et al. Specific plant terpenoids and lignoids possess potent antiviral activities against severe acute respiratory syndrome coronavirus. *J Med Chem* 2007;**50**:4087–95.
- 146. Zahedipour F, Hosseini SA, Sathyapalan T, Majeed M, Jamialahmadi T, Al-Rasadi K, et al. Potential effects of curcumin in the treatment of COVID-19 infection. *Phytother Res* 2020;**34**:2911–20.
- 147. Nguyen TT, Woo HJ, Kang HK, Nguyen VD, Kim YM, Kim DW, et al. Flavonoid-mediated inhibition of SARS coronavirus 3C-like protease expressed in Pichia pastoris. *Biotechnol Lett* 2012;**34**:831–8.

- 148. Abian O, Ortega-Alarcon D, Jimenez-Alesanco A, Ceballos-Laita L, Vega S, Reyburn HT, et al. Structural stability of SARS-CoV-2 3CLpro and identification of quercetin as an inhibitor by experimental screening. *Int J Biol Macromol* 2020;**164**:1693–703.
- 149. Park JY, Kim JH, Kim YM, Jeong HJ, Kim DW, Park KH, et al. Tanshinones as selective and slow-binding inhibitors for SARS-CoV cysteine proteases. *Bioorg Med Chem* 2012;20:5928–35.
- 150. Park JY, Ko JA, Kim DW, Kim YM, Kwon HJ, Jeong HJ, et al. Chalcones isolated from *Angelica keiskei* inhibit cysteine proteases of SARS-CoV. *J Enzyme Inhib Med Chem* 2016;**31**:23–30.
- 151. Lin CW, Tsai FJ, Tsai CH, Lai CC, Wan L, Ho TY, et al. Anti-SARS coronavirus 3C-like protease effects of Isatis indigotica root and plant-derived phenolic compounds. *Antiviral Res* 2005;**68**:36–42.
- 152. Jo S, Kim S, Shin DH, Kim MS. Inhibition of SARS-CoV 3CL protease by flavonoids. *J Enzyme Inhib Med Chem* 2020;**35**:145–51.
- 153. Yi L, Li Z, Yuan K, Qu X, Chen J, Wang G, et al. Small molecules blocking the entry of severe acute respiratory syndrome coronavirus into host cells. *J Virol* 2004;78:11334–9.
- 154. Ryu YB, Jeong HJ, Kim JH, Kim YM, Park JY, Kim D, et al. Biflavonoids from Torreya nucifera displaying SARS-CoV 3CL(pro) inhibition. Bioorg Med Chem 2010;18:7940–7.
- 155. Park JY, Jeong HJ, Kim JH, Kim YM, Park SJ, Kim D, et al. Diarylheptanoids from *Alnus japonica* inhibit papain-like protease of severe acute respiratory syndrome coronavirus. *Biol Pharm Bull* 2012;**35**:2036–42.
- 156. Song YH, Kim DW, Curtis-Long MJ, Yuk HJ, WangY, Zhuang N, et al. Papain-like protease (PLpro) inhibitory effects of cinnamic amides from *Tribulus* terrestris fruits. *Biol Pharm Bull* 2014;**37**:1021–8.
- 157. Kim DW, Seo KH, Curtis-Long MJ, Oh KY, Oh JW, Cho JK, et al. Phenolic

- phytochemical displaying SARS-CoV papain-like protease inhibition from the seeds of *Psoralea corylifolia*. *J Enzyme Inhib Med Chem* 2014;**29**:59–63.
- 158. Cho JK, Curtis-Long MJ, Lee KH, Kim DW, Ryu HW, Yuk HJ, et al. Geranylated flavonoids displaying SARS-CoV papain-like protease inhibition from the fruits of *Paulownia tomentosa*. *Bioorg Med Chem* 2013;**21**:3051–7.
- 159. Cinatl J, Morgenstern B, Bauer G, Chandra P, Rabenau H, Doerr HW. Glycyrrhizin, an active component of liquorice roots, and replication of SARS-associated coronavirus. *Lancet* 2003;**361**:2045–6.
- 160. Hoever G, Baltina L, Michaelis M, Kondratenko R, Baltina L, Tolstikov GA, et al. Antiviral activity of glycyrrhizic acid derivatives against SARS-coronavirus. J Med Chem 2005;48:1256–9.
- 161. Bailly C, Vergoten G. Glycyrrhizin: an alternative drug for the treatment of COVID-19 infection and the associated respiratory syndrome? *Pharmacol Ther* 2020;**214**:107618.
- 162. Zhang CH, Wang YF, Liu XJ, Lu JH, Qian CW, Wan ZY, et al. Antiviral activity of cepharanthine against severe acute respiratory syndrome coronavirus *in vitro*. *Chin Med J (Engl)* 2005;**118**:493–6.
- 163. Wu CY, Jan JT, Ma SH, Kuo CJ, Juan HF, Cheng YS, et al. Small molecules targeting severe acute respiratory syndrome human coronavirus. *Proc Natl Acad Sci U S A* 2004;**101**:10012–7.
- 164. Li SY, Chen C, Zhang HQ, Guo HY, Wang H, Wang L, et al. Identification of natural compounds with antiviral activities against SARS-associated coronavirus. *Antiviral Res* 2005;67:18–23.
- 165. Bridgford JL, Xie SC, Cobbold SA, Pasaje CFA, Herrmann S, Yang T, et al. Artemisinin kills malaria parasites by damaging proteins and inhibiting the proteasome. *Nat Commun* 2018;**9**:3801.
- 166. Wang J, Zhang J, Shi Y, Xu C, Zhang C, Wong YK, et al. Mechanistic investigation of the specific anticancer property of artemisinin and its

- combination with aminolevulinic acid for enhanced anticolorectal cancer activity. *ACS Cent Sci* 2017;**3**:743–50.
- 167. An J, Minie M, Sasaki T, Woodward JJ, Elkon KB. Antimalarial drugs as immune modulators: new mechanisms for old drugs. *Annu Rev Med* 2017;**68**:317–30.
- 168. Huang F, Li Y, Leung EL, Liu X, Liu K, Wang Q, et al. A review of therapeutic agents and Chinese herbal medicines against SARS-COV-2 (COVID-19). *Pharmacol Res* 2020;**158**:104929.
- 169. Cao B, Hayden FG. Antiviral monotherapy for hospitalised patients with COVID-19 is not enough. *Lancet* 2020;**396**:1310–1.
- 170. Huang CF, Lin SS, Liao PH, Young SC, Yang CC. The immunopharmaceutical effects and mechanisms of herb medicine. *Cell Mol Immunol* 2008;**5**:23–31.
- 171. Huang YF, Bai C, He F, Xie Y, Zhou H. Review on the potential action mechanisms of Chinese medicines in treating Coronavirus Disease 2019 (COVID-19). *Pharmacol Res* 2020;**158**:104939.
- 172. Ma HD, Deng YR, Tian Z, Lian ZX. Traditional Chinese medicine and immune regulation. *Clin Rev Allergy Immunol* 2013;44:229–41.
- 173. Cao ZY, Liu YZ, Li JM, Ruan YM, Yan WJ, Zhong SY, et al. Glycyrrhizic acid as an adjunctive treatment for depression through anti-inflammation: a randomized placebo-controlled clinical trial. *J Affect Disord* 2020;**265**:247–54.
- 174. Zhang C, Wu Z, Li JW, Zhao H, Wang GQ. Cytokine release syndrome in severe COVID-19: interleukin-6 receptor antagonist tocilizumab may be the key to reduce mortality. *Int J Antimicrob Agents* 2020;**55**:105954.
- 175. Crisafulli S, Isgro V, La Corte L, Atzeni F, Trifiro G. Potential role of antiinterleukin (IL)-6 drugs in the treatment of COVID-19: rationale, clinical evidence and risks. *BioDrugs* 2020;**34**:415–22.
- 176. Chakraborty C, Sharma AR, Bhattacharya M, Sharma G, Lee SS, Agoramoorthy G. COVID-19: consider IL-6 receptor antagonist for the therapy of cytokine

- storm syndrome in SARS-CoV-2 infected patients. *J Med Virol* 2020;**92**:2260–2.
- 177. Mehta P, McAuley DF, Brown M, Sanchez E, Tattersall RS, Manson JJ, et al. COVID-19: consider cytokine storm syndromes and immunosuppression. *Lancet* 2020;**395**:1033–4.
- 178. Moore JB, June CH. Cytokine release syndrome in severe COVID-19. *Science* 2020;**368**:473–4.
- 179. Xu X, Han M, Li T, Sun W, Wang D, Fu B, et al. Effective treatment of severe COVID-19 patients with tocilizumab. *Proc Natl Acad Sci U S A* 2020;**117**:10970–5.
- 180. Masia M, Fernandez-Gonzalez M, Padilla S, Ortega P, Garcia JA, Agullo V, et al. Impact of interleukin-6 blockade with tocilizumab on SARS-CoV-2 viral kinetics and antibody responses in patients with COVID-19: a prospective cohort study. *EBioMedicine* 2020;60:102999.
- 181. Dzobo K, Chiririwa H, Dandara C, Dzobo W. Coronavirus disease-2019 treatment strategies targeting interleukin-6 signaling and herbal medicine. *OMICS* 2020;**25**:13–22.
- 182. Nasonov E, Samsonov M. The role of Interleukin 6 inhibitors in therapy of severe COVID-19. *Biomed Pharmacother* 2020;**131**:110698.
- 183. Tang Y, Liu J, Zhang D, Xu Z, Ji J, Wen C. Cytokine storm in COVID-19: the current evidence and treatment strategies. *Front Immunol* 2020;**11**:1708.
- 184. Song Y, Yao C, Shang H, Yao X, Bai C. Intravenous infusion of Chinese medicine Xuebijing for patients with severe pneumonia: a multicenter, randomised, double-blind controlled trial. *The Lancet* 2017;**390**:S34.
- 185. Lyu M, Zhou Z, Wang X, Lv H, Wang M, Pan G, et al. Network pharmacology-guided development of a novel integrative regimen to prevent acute graft-vs.-host disease. *Front Pharmacol* 2018;**9**:1440.
- 186. Wang L, Liu Z, Dong Z, Pan J, Ma X. Effects of Xuebijing injection on microcirculation in septic shock. *J Surg Res* 2016;**202**:147–54.

- 187. Yin Q, Li C. Treatment effects of xuebijing injection in severe septic patients with disseminated intravascular coagulation. *Evid Based Complement Alternat Med* 2014;**2014**:949254.
- 188. Song Y, Yao C, Yao Y, Han H, Zhao X, Yu K, et al. XueBiJing injection *versus* placebo for critically ill patients with severe community-acquired pneumonia: a randomized controlled trial. *Crit Care Med* 2019;47:e735–43.
- 189. Hou SY, Feng XH, Lin CL, Tan YF. Efficacy of Xuebijing for coagulopathy in patients with sepsis. *Saudi Med J* 2015;**36**:164–9.
- 190. Liu MW, Su MX, Zhang W, Wang YQ, Chen M, Wang L, et al. Protective effect of Xuebijing injection on paraquat-induced pulmonary injury *via* down-regulating the expression of p38 MAPK in rats. *BMC Complement Altern Med* 2014;**14**:498.
- 191. He F, Wang J, Liu Y, Wang X, Cai N, Wu C, et al. Xuebijing injection induces anti-inflammatory-like effects and downregulates the expression of TLR4 and NF-kappaB in lung injury caused by dichlorvos poisoning. *Biomed Pharmacother* 2018;**106**:1404–11.
- 192. Chen X, Feng Y, Shen X, Pan G, Fan G, Gao X, et al. Anti-sepsis protection of Xuebijing injection is mediated by differential regulation of pro- and anti-inflammatory Th17 and T regulatory cells in a murine model of polymicrobial sepsis. *J Ethnopharmacol* 2018;**211**:358–65.
- 193. Liu MW, Wang YH, Qian CY, Li H. Xuebijing exerts protective effects on lung permeability leakage and lung injury by upregulating Toll-interacting protein expression in rats with sepsis. *Int J Mol Med* 2014;**34**:1492–504.
- 194. Zhang H, Wei L, Zhao G, Liu S, Zhang Z, Zhang J, et al. Protective effect of Xuebijing injection on myocardial injury in patients with sepsis: a randomized clinical trial. *J Tradit Chin Med* 2016;**36**:706–10.
- 195. Yuxi Q, Zhang H, Baili Y, Shi S. Effects of Xuebijing injection for patients with sepsis-induced acute kidney injury after Wenchuan earthquake. *Altern Ther*

- Health Med 2017;23:36-42.
- 196. Ji M, Wang Y, Wang L, Chen L, Li J. Protective effect of Xuebijing injection against acute lung injury induced by left ventricular ischemia/reperfusion in rabbits. *Exp Ther Med* 2016;**12**:51–8.
- 197. Xu Q, Liu J, Guo X, Tang Y, Zhou G, Liu Y, et al. Xuebijing injection reduces organ injuries and improves survival by attenuating inflammatory responses and endothelial injury in heatstroke mice. *BMC Complement Altern Med* 2015;**15**:4.
- 198. Liu X, Hu Z, Zhou B, Li X, Tao R. Chinese herbal preparation Xuebijing potently inhibits inflammasome activation in hepatocytes and ameliorates mouse liver ischemia–reperfusion injury. *PLoS One* 2015;**10**:e0131436.
- 199. Chen Y, Tong H, Zhang X, Tang L, Pan Z, Liu Z, et al. Xuebijing injection alleviates liver injury by inhibiting secretory function of Kupffer cells in heat stroke rats. *J Tradit Chin Med* 2013;**33**:243–9.
- 200. He YQ, Zhou CC, Yu LY, Wang L, Deng JL, Tao YL, et al. Natural product derived phytochemicals in managing acute lung injury by multiple mechanisms. *Pharmacol Res* 2020;**163**:105224.
- 201. Dong S, Zhong Y, Yang K, Xiong X, Mao B. Intervention effect and dose-dependent response of tanreqing injection on airway inflammation in lipopolysaccharide-induced rats. *J Tradit Chin Med* 2013;33:505–12.
- 202. Zhong Y, Mao B, Wang G, Fan T, Liu X, Diao X, et al. Tanreqing injection combined with conventional Western medicine for acute exacerbations of chronic obstructive pulmonary disease: a systematic review. *J Altern Complement Med* 2010;**16**:1309–19.
- 203. Liu W, Jiang HL, Cai LL, Yan M, Dong SJ, Mao B. Tanreqing injection attenuates lipopolysaccharide-induced airway inflammation through MAPK/NF-kappaB signaling pathways in rats model. *Evid Based Complement Alternat Med* 2016;**2016**:5292346.
- 204. Liu W, Zhang X, Mao B, Jiang H. Systems pharmacology-based study of

- Tanreqing injection in airway mucus hypersecretion. *J Ethnopharmacol* 2020;**249**:112425.
- 205. Jiang C, Zhong R, Zhang J, Wang X, Ding G, Xiao W, et al. Reduning injection ameliorates paraquat-induced acute lung injury by regulating AMPK/MAPK/NF-kappaB signaling. *J Cell Biochem* 2019;**120**:12713–23.
- 206. Chen W, Ma Y, Zhang H, Guo Y, Guan M, Wang Y. Reduning plus ribavirin display synergistic activity against severe pneumonia induced by H1N1 influenza A virus in mice. *J Tradit Chin Med* 2020;**40**:803–11.
- 207. Tang LP, Xiao W, Li YF, Li HB, Wang ZZ, Yao XS, et al. Anti-inflammatory effects of Reduning Injection on lipopolysaccharide-induced acute lung injury of rats. *Chin J Integr Med* 2014;**20**:591–9.
- 208. Peng S, Hang N, Liu W, Guo W, Jiang C, Yang X, et al. Andrographolide sulfonate ameliorates lipopolysaccharide-induced acute lung injury in mice by down-regulating MAPK and NF-kappaB pathways. *Acta Pharm Sin B* 2016;**6**:205–11.
- 209. Shi H, Guo W, Zhu H, Li M, Ung COL, Hu H, et al. Cost-effectiveness analysis of Xiyanping injection (andrographolide sulfonate) for treatment of adult community acquired pneumonia: a retrospective, propensity score-matched cohort study. *Evid Based Complement Alternat Med* 2019;**2019**:4510591.
- 210. Liu W, Guo W, Guo L, Gu Y, Cai P, Xie N, et al. Andrographolide sulfonate ameliorates experimental colitis in mice by inhibiting Th1/Th17 response. *Int Immunopharmacol* 2014;**20**:337–45.
- 211. Yin Q, Liu B, Wu C, Yang J, Hang C, Li C. Effects of Shen-Fu injection on coagulation-fibrinolysis disorders in a porcine model of cardiac arrest. Am J Emerg Med 2016;34:469–76.
- 212. Yan X, Wu H, Ren J, Liu Y, Wang S, Yang J, et al. Shenfu Formula reduces cardiomyocyte apoptosis in heart failure rats by regulating microRNAs. *J Ethnopharmacol* 2018;**227**:105–12.

- 213. Mao ZJ, Zhang Q L, Shang J, Gao T, Yuan WJ, Qin LP. Shenfu Injection attenuates rat myocardial hypertrophy by up-regulating miR-19a-3p expression. *Sci Rep* 2018;**8**:4660.
- 214. Zhang Q, Li C, Shao F, Zhao L, Wang M, Fang Y. Efficacy and safety of combination therapy of Shenfu injection and postresuscitation bundle in patients with return of spontaneous circulation after in-hospital cardiac arrest: a randomized, assessor-blinded, controlled trial. *Crit Care Med* 2017;45:1587–95.
- 215. Jin YY, Gao H, Zhang XY, Ai H, Zhu XL, Wang J. Shenfu Injection inhibits inflammation in patients with acute myocardial infarction complicated by cardiac shock. *Chin J Integr Med* 2017;**23**:170–5.
- 216. Wang YY, Li YY, Li L, Yang DL, Zhou K, Li YH. Protective effects of Shenfu injection against myocardial ischemia–reperfusion injury *via* activation of eNOS in rats. *Biol Pharm Bull* 2018;**41**:1406–13.
- 217. Shao F, Li H, Li D, Li C. Effects of Shenfu injection on survival and neurological outcome after out-of-hospital cardiac arrest: a randomised controlled trial. *Resuscitation* 2020;**150**:139–44.
- 218. Wu J, Li C, Yuan W. Effects of Shenfu injection on macrocirculation and microcirculation during cardiopulmonary resuscitation. *J Ethnopharmacol* 2016;**180**:97–103.
- 219. Ni J, Shi Y, Li L, Chen J, Li L, Li M, et al. Cardioprotection against heart failure by Shenfu Injection *via* TGF-beta/Smads signaling pathway. *Evid Based Complement Alternat Med* 2017;**2017**:7083016.
- 220. Tan G, Zhou Q, Liu K, Dong X, Li L, Liao W, et al. Cross-platform metabolic profiling deciphering the potential targets of Shenfu injection against acute viral myocarditis in mice. *J Pharm Biomed Anal* 2018;**160**:1–11.
- 221. Xu P, Zhang W Q, Xie J, Wen YS, Zhang GX, Lu SQ. Shenfu injection prevents sepsis-induced myocardial injury by inhibiting mitochondrial apoptosis. *J Ethnopharmacol* 2020;**261**:113068.

- 222. Zhang XJ, Song L, Zhou ZG, Wang XM. Effect of shenfu injection on gastrointestinal microcirculation in rabbits after myocardial ischemia—reperfusion injury. *World J Gastroenterol* 2006;**12**:4389–91.
- 223. Jin S, Jiang R, Lei S, Jin L, Zhu C, Feng W, et al. Shenfu injection prolongs survival and protects the intestinal mucosa in rats with sepsis by modulating immune response. *Turk J Gastroenterol* 2019;**30**:364–71.
- 224. Ma X, Yang YX, Chen N, Xie Q, Wang T, He X, et al. Meta-analysis for clinical evaluation of Xingnaojing injection for the treatment of cerebral infarction. *Front Pharmacol* 2017;**8**:485.
- 225. Zhang YM, Qu XY, Tao LN, Zhai JH, Gao H, Song YQ, et al. XingNaoJing injection ameliorates cerebral ischaemia/reperfusion injury *via* SIRT1-mediated inflammatory response inhibition. *Pharm Biol* 2020;**58**:16–24.
- 226. Xu M, Su W, Xu QP, Huang WD. Effect of Xingnaojing injection on cerebral edema and blood–brain barrier in rats following traumatic brain injury. *Chin J Traumatol* 2010;**13**:158–62.
- 227. Qu XY, Zhang YM, Tao LN, Gao H, Zhai JH, Sun JM, et al. XingNaoJing injections protect against cerebral ischemia/reperfusion injury and alleviate blood–brain barrier disruption in rats, through an underlying mechanism of NLRP3 inflammasomes suppression. *Chin J Nat Med* 2019;**17**:498–505.
- 228. Ma X, Wang T, Wen J, Wang J, Zeng N, Zou W, et al. Role of Xingnaojing injection in treating acute cerebral hemorrhage: a systematic review and meta-analysis. *Medicine (Baltimore)* 2020;**99**:e19648.
- 229. Liu H, Yan Y, Pang P, Mao J, Hu X, Li D, et al. Angong Niuhuang Pill as adjuvant therapy for treating acute cerebral infarction and intracerebral hemorrhage: a meta-analysis of randomized controlled trials. *J Ethnopharmacol* 2019;**237**:307–13.
- 230. Zhang DS, Liu YL, Zhu DQ, Huang XJ, Luo CH. Point application with Angong Niuhuang sticker protects hippocampal and cortical neurons in rats with cerebral

- ischemia. Neural Regen Res 2015;10:286–91.
- 231. Zhang D, Fu M, Song C, Wang C, Lin X, Liu Y. Expressions of apoptosis-related proteins in rats with focal cerebral ischemia after Angong Niuhuang sticker point application. *Neural Regen Res* 2012;7:2347–53.
- 232. Huang K, Zhang P, Zhang Z, Youn JY, Zhang H, Cai HL. Traditional Chinese medicine (TCM) in the treatment of viral infections: efficacies and mechanisms. *Pharmacol Ther* 2021;**225**:107843.
- 233. Yeoh YK, Zuo T, Lui GC, Zhang F, Liu Q, Li AY, et al. Gut microbiota composition reflects disease severity and dysfunctional immune responses in patients with COVID-19. *Gut* 2021;**70**:698–706.
- 234. Augusti PR, Conterato GMM, Denardin CC, Prazeres ID, Serra AT, Bronze MR, et al. Bioactivity, bioavailability, and gut microbiota transformations of dietary phenolic compounds: implications for COVID-19. *J Nutr Biochem* 2021;97:108787.
- 235. Li X, Wu D, Niu J, Sun Y, Wang Q, Yang B, et al. Intestinal flora: a pivotal role in investigation of traditional Chinese medicine. *Am J Chin Med* 2021;49:237–68.
- 236. Zhang Q, Yue S, Wang W, Chen Y, Zhao C, Song Y, et al. Potential role of gut microbiota in traditional Chinese medicine against COVID-19. *Am J Chin Med* 2021;49:785–803.
- 237. Wu GS, Zhong J, Zheng NN, Wang CR, Jin HL, Ge GB, et al. Investigation of modulating effect of Qingfei Paidu Decoction on host metabolism and gut microbiome in rats. *Chin J Chin Mater Med* 2020;**45**:3726–39.
- 238. Lyu M, Wang YF, Fan GW, Wang XY, Xu SY, Zhu Y. Balancing herbal medicine and functional food for prevention and treatment of cardiometabolic diseases through modulating gut microbiota. *Front Microbiol* 2017;8:2146.
- 239. Qiu Y, Yang J, Wang L, Yang X, Gao K, Zhu C, et al. Dietary resveratrol attenuation of intestinal inflammation and oxidative damage is linked to the

- alteration of gut microbiota and butyrate in piglets challenged with deoxynivalenol. *J Anim Sci Biotechnol* 2021;**12**:71.
- 240. Valizadeh H, Abdolmohammadi-Vahid S, Danshina S, Ziya GM, Ammari A, Sadeghi A, et al. Nano-curcumin therapy, a promising method in modulating inflammatory cytokines in COVID-19 patients. Int Immunopharmacol 2020;89:107088.
- 241. Dai YJ, Wan SY, Gong SS, Liu JC, Li F, Kou JP. Recent advances of traditional Chinese medicine on the prevention and treatment of COVID-19. *Chin J Nat Med* 2020;**18**:881–9.
- 242. Wang Z, Yang L. Chinese herbal medicine: fighting SARS-CoV-2 infection on all fronts. *J Ethnopharmacol* 2021;**270**:113869.
- 243. Lee DYW, Li QY, Liu J, Efferth T. Traditional Chinese herbal medicine at the forefront battle against COVID-19: clinical experience and scientific basis. *Phytomedicine* 2020;**80**:153337.
- 244. An X, Zhang Y, Duan L, Jin, Zhao S, Zhou R, et al. The direct evidence and mechanism of traditional Chinese medicine treatment of COVID-19. *Biomed Pharmacother* 2021;**137**:111267.
- 245. Huang J, Tao G, Liu J, Cai J, Huang Z, Chen JX. Current prevention of COVID-19: natural products and herbal medicine. *Front Pharmacol* 2020;**11**:588508.
- 246. Jalali A, Dabaghian F, Akbrialiabad H, Foroughinia F, Zarshenas MM. A pharmacology-based comprehensive review on medicinal plants and phytoactive constituents possibly effective in the management of COVID-19. *Phytother Res* 2020;**35**:1925–38.
- 247. Nugraha RV, Ridwansyah H, Ghozali M, Khairani AF, Atik N. Traditional herbal medicine candidates as complementary treatments for COVID-19: a review of their mechanisms, pros and cons. *Evid Based Complement Alternat Med* 2020;**2020**:2560645.
- 248. Li Q, Ran Q, Sun L, Yin J, Luo T, Liu L, et al. Lian Hua Qing Wen capsules, a

- potent epithelial protector in acute lung injury model, block proapoptotic communication between macrophages, and alveolar epithelial cells. *Front Pharmacol* 2020;**11**:522729.
- 249. Chen K, Chen H. Traditional Chinese medicine for combating COVID-19. *Front Med* 2020;**14**:529–32.
- 250. Huang ST, Lai HC, Lin YC, Huang WT, Hung HH, Ou SC, et al. Principles and treatment strategies for the use of Chinese herbal medicine in patients at different stages of coronavirus infection. *Am J Cancer Res* 2020;**10**:2010–31.
- 251. Yang Y, Islam MS, Wang J, Li Y, Chen X. Traditional Chinese medicine in the treatment of patients infected with 2019-new coronavirus (SARS-CoV-2): a review and perspective. *Int J Biol Sci* 2020;**16**:1708–17.
- 252. Khare P, Sahu U, Pandey SC, Samant M. Current approaches for target-specific drug discovery using natural compounds against SARS-CoV-2 infection. *Virus Res* 2020;290:198169.
- 253. Wiersinga WJ, Rhodes A, Cheng AC, Peacock SJ, Prescott HC. Pathophysiology, transmission, diagnosis, and treatment of coronavirus disease 2019 (COVID-19): a review. *JAMA* 2020;**324**:782–93.
- 254. Dhama K, Khan S, Tiwari R, Sircar S, Bhat S, Malik YS, et al. Coronavirus disease 2019-COVID-19. *Clin Microbiol Rev* 2020;**33**:e00028-20.
- 255. Xian Y, Zhang J, Bian Z, Zhou H, Zhang Z, Lin Z, et al. Bioactive natural compounds against human coronaviruses: a review and perspective. *Acta Pharm Sin B* 2020;**10**:1163–74.
- 256. Li S, Liu C, Guo F, Taleb SJ, Tong M, Shang D. Traditional Chinese medicine as potential therapy for COVID-19. *Am J Chin Med* 2020;**48**:1263–77.
- 257. Shi J, Lu Y, Zhang Y, Xia L, Ye C, Lu Y, et al. Traditional Chinese medicine formulation therapy in the treatment of coronavirus disease 2019 (COVID-19). *Am J Chin Med* 2020;**48**:1523–38.
- 258. Zhang L, Yu J, Zhou Y, Shen M, Sun L. Becoming a faithful defender: traditional

- Chinese medicine against coronavirus disease 2019 (COVID-19). *Am J Chin Med* 2020;**48**:763–77.
- 259. Sun CY, Sun YL, Li XM. The role of Chinese medicine in COVID-19 pneumonia: a systematic review and meta-analysis. *Am J Emerg Med* 2020;**38**:2153–9.
- 260. Zhang J, Xie B, Hashimoto K. Current status of potential therapeutic candidates for the COVID-19 crisis. *Brain Behav Immun* 2020;**87**:59–73.
- 261. Liu L. Traditional Chinese medicine contributes to the treatment of COVID-19 patients. *Chin Herb Med* 2020;**12**:95–6.
- 262. Li JG, Xu H. Chinese medicine in fighting against Covid-19: role and inspiration. *Chin J Integr Med* 2021;**27**:3–6.
- 263. Tong T, Wu YQ, Ni WJ, Shen AZ, Liu S. The potential insights of traditional Chinese medicine on treatment of COVID-19. *Chin Med* 2020;**15**:51.
- 264. Xu J, Zhang Y. Traditional Chinese medicine treatment of COVID-19. Complement Ther Clin Pract 2020;39:101165.
- 265. Zhuang W, Fan Z, Chu Y, Wang H, Yang Y, Wu L, et al. Chinese patent medicines in the treatment of coronavirus disease 2019 (COVID-19) in China. *Front Pharmacol* 2020;**11**:1066.
- 266. Lopez-Alcalde J, Yan Y, Witt C M, Barth J. Current state of research about Chinese herbal medicines (CHM) for the treatment of coronavirus disease 2019 (COVID-19): a scoping review. *J Altern Complement Med* 2020;**26**:557–70.
- 267. Zhou LP, Wang J, Xie RH, Pakhale S, Krewski D, Cameron DW, et al. The effects of traditional Chinese medicine as an auxiliary treatment for COVID-19: a systematic review and meta-analysis. *J Altern Complement Med* 2021;**27**:225–37.
- 268. Ling CQ. Traditional Chinese medicine is a resource for drug discovery against 2019 novel coronavirus (SARS-CoV-2). *J Integr Med* 2020;**18**:87–8.
- 269. Al-Romaima A, Liao Y, Feng J, Qin X, Qin G. Advances in the treatment of

- novel coronavirus disease (COVID-19) with Western medicine and traditional Chinese medicine: a narrative review. *J Thorac Dis* 2020;**12**:6054–69.
- 270. McKee DL, Sternberg A, Stange U, Laufer S, Naujokat C. Candidate drugs against SARS-CoV-2 and COVID-19. *Pharmacol Res* 2020;**157**:104859.
- 271. Ren JL, Zhang AH, Wang XJ. Traditional Chinese medicine for COVID-19 treatment. *Pharmacol Res* 2020;**155**:104743.
- 272. Wang WY, Xie Y, Zhou H, Liu L. Contribution of traditional Chinese medicine to the treatment of COVID-19. *Phytomedicine* 2021;**85**:153279.
- 273. Zhang JL, Li WX, Li Y, Wong MS, Wang YJ, Zhang Y. Therapeutic options of TCM for organ injuries associated with COVID-19 and the underlying mechanism. *Phytomedicine* 2021;**85**:153297.
- 274. Zhao Z, Li Y, Zhou L, Zhou X, Xie B, Zhang W, et al. Prevention and treatment of COVID-19 using traditional Chinese medicine: a review. *Phytomedicine* 2021;85:153308.
- 275. He T, Qu R, Qin C, Wang Z, Zhang Y, Shao X, et al. Potential mechanisms of Chinese herbal medicine that implicated in the treatment of COVID-19 related renal injury. *Saudi Pharm J* 2020;**28**:1138–48.
- 276. Cui HT, Li YT, Guo LY, Liu XG, Wang LS, Jia JW, et al. Traditional Chinese medicine for treatment of coronavirus disease 2019: a review. *Trad Med Res* 2020;**5**:65–73.
- 277. Mani JS, Johnson JB, Steel JC, Broszczak DA, Neilsen PM, Walsh KB, et al. Natural product-derived phytochemicals as potential agents against coronaviruses: a review. *Virus Res* 2020;284:197989.
- 278. Gray PE, Belessis Y. The use of Traditional Chinese medicines to treat SARS-CoV-2 may cause more harm than good. *Pharmacol Res* 2020;**156**:104776.
- 279. Yang Y. Use of herbal drugs to treat COVID-19 should be with caution. *Lancet* 2020;**395**:1689–90.
- 280. Zhang AH, Ren JL, Wang XJ. Reply to "The use of traditional Chinese

- medicines to treat SARS-CoV-2 may cause more harm than good". *Pharmacol Res* 2020;**157**:104775.
- 281. Huang C, Huang L, Wang Y, Li X, Ren L, Gu X, et al. 6-Month consequences of COVID-19 in patients discharged from hospital: a cohort study. *Lancet* 2021;**397**:220–32.
- 282. Yelin D, Wirtheim E, Vetter P, Kalil AC, Bruchfeld J, Runold M, et al. Long-term consequences of COVID-19: research needs. *Lancet Infect Dis* 2020;**20**:1115–7.
- 283. An YW, Yuan B, Wang JC, Wang C, Liu TT, Song S, et al. Clinical characteristics and impacts of traditional Chinese medicine treatment on the convalescents of COVID-19. *Int J Med Sci* 2021;**18**:646–51.

Figure captions

Figure 1 The recommended Chinese patent medicines or Chinese medicine formulas for distinct stages of COVID-19 treatment.

Figure 2 Representative herbs and their main active ingredients and functions for COVID-19. (A) The herb-ingredient-target-function network of frequently used herbs and their main ingredients, as well as their key targets and functions for COVID-19. (B) The chemical structures of main active ingredients and their main functions of commonly used herbs for COVID-19.

Figure 3 An overview of pathogenesis of COVID-19 and the potential mechanisms of TCM remedy in distinct disease stages.

Table 1 Clinical efficacies of integrated TCM and WM for COVID-19 treatment.

No.	Intervention	Method	Object	Disease stage	Clinical manifestation	Laboratory finding	Ref.
			(T/C)				
1	Jinhua Qinggan granules+WM vs.	Retrospective	44/36	Moderate or	1) Shorten the duration of nucleic acid turn negative	Increase WBC and	36
	WM	cohort study		severe	2) Promote the absorption of pneumonia inflammatory	lymphocyte count	
		(RCS)			exudate		
2	Jinhua Qinggan capsule+WM vs.	Randomized	82/41	Mild	Reduce the symptoms of fever, cough, fatigue, and	Unreported	37
	WM	controlled trial			sputum cough, and relieve the psychological anxiety		
		(RCT)					
3	Lianhua Qingwen capsule+WM	RCT	142/142	Mild or	1) Shorten median time to symptom recovery	Unreported	27
	vs. WM			moderate	2) Shorten time to recovery of fever, fatigue, and cough		
					3) Improve the rate of chest CT manifestations and		
					clinical cure		
4	Lianhua Qingwen capsule+WM	RCS	63/38	All	Relieve symptoms of fever, cough, weakness, and short	Unreported	38
	vs. WM				of breath		
5	Lianhua Qingwen	RCT	147/148	Mild or	Relieve symptoms of fever, fatigue, cough, dry throat,	Lower the levels of CRP	39
	capsule+arbidol vs. arbidol			moderate	sore throat, and chest tightness	and procalcitonin,	
						elevate WBC and	
						lymphocyte count	
6	Lianhua Qingwen capsule+WM	Before and	54/0	Moderate	Relieve the symptoms and reduce the duration of fever,	Unreported	40
		after			fatigue, and cough.		
		comparison					
7	Lianhua Qingwen	RCT	189/94	All	1) Improve the symptoms of fever and diarrhea,	Unreported	28
	capsule+Huoxiang Zhengqi				especially fatigue, nausea and vomiting, chest tightness,		
	dropping pills+WM vs. WM				shortness of breath and limb soreness		
					2) Reduce the utilization rate of anti-infective drugs and		
					improve the prognosis of patients		
					3) Block disease aggravation		
8	Lianhua Qingwen	RCS	68/40	Mild or	1) Shorten the median time from admission to the first	1) Increase lymphocytes	41

	capsule+arbidol vs. arbidol			moderate	negative result of nucleic acid detection	count	
					2) Reduce lung inflammation	2) Lower the levels of	
						serum amyloid A and	
						CRP	
9	Xuebijing injection+WM	Case analysis	11/0	Severe or	May ameliorate lung injury	Reduce the levels of	42
				critical		TNF- α , IP-10, MIP-1 β ,	
						and RANTES	
10	Xuebijing injection+WM vs. WM	RCT	40/20	Severe	Improve the conditions of patients, lower APACHE II	1) Improve the	43
					score	oxygenation index of	
						PaO ₂ /FiO ₂	
						2) Increase WBC and	
						lymphocyte count,	
						decrease the levels of	
						CRP and ESR	
11	Xuebijing injection+antiviral	RCS	22/22	Moderate	Increase the effective rate of lung lesions absorption	Tend to improve WBC	44
	treatment vs. antiviral treatment				and the overall effective rate of treatment	count, lymphocyte	
						count, and the levels of	
						CRP and ferritin	
12	Qingfei Paidu decoction+WM vs.	RCS	37/26	Severe	1) Relieve the symptoms and improve inflammation	1) Improve the levels of	45
	WM				resolution in the lung	CRP, CK, creatine	
					2) Tend to mitigate the extent of multi-organ	kinase-myocardial band,	
					impairment	LDH, and blood urea	
						nitrogen	
						2) Increase lymphocyte	
						count	
13	Qingfei Paidu decoction+WM	Before and	98/0	All	1) Nearly all adverse symptoms including fever, cough,	Restore the levels of	46
		after			asthma, and fatigue were relieved	AST, ALT, D-dimer,	
		comparison			2) Improve lung CT imaging	CRP, ESR, and the	
						percentage of	

						lymphocyte	
14	Qingfei Paidu decoction+antiviral	RCS	30/30	All	1) Shorten inpatient days and reduce the time of fever	Unreported	47
	treatment vs. antiviral treatment				and cough		
					2) Promote lung CT improvement		
15	Qingfei Paidu decoction+WM	RCS	46/43	All	1) Reduce inflammation, enhance cellular immunity,	Reduce the level of IL-6	48
					improve renal function, lower hypercoagulability	and increase the level of	
					2) Shorten the length of hospitalization and nucleic acid	CD3	
					negative time		
16	Qingfei Paidu decoction+WM vs.	Multi-center	199/96	Mild or	1) Reduce mean length of hospital stay, nucleic acid	Unreported	49
	WM	clinical		moderate	negative time and improve symptom of sputum		
		observation			2) Improve lung CT imaging		
17	Qingfei Paidu decoction+WM	Multi-center	782/0	All	Shorten the time of recovery, viral shedding, and the	Unreported	50
		clinical			duration of hospital stay		
		observation					
18	Xuanfei Baidu Decoction+WM	RCT	22/20	Mild	Increase the disappearance rate of symptoms of fever,	1) Elevate WBC and	32
	vs. WM				cough, fatigue, and loss of appetite	lymphocyte count	
						2) Reduce the levels of	
						CRP and ESR	
19	Huashi Baidu granule vs. WM	RCS	23/32	Severe	Improve chest CT imaging and lung lesion opacity	Decrease the levels of	30
						CRP, ESR, serum	
						ferritin, and myoglobin	
20	Huashi Baidu formula+TCM	RCS	20/20/20	Mild or	Shorten the clinical remission time	No significant	31
	injection vs. Huashi Baidu			moderate		differences in	
	formula+lopinavir–ritonavir vs.					biochemical indicators	
	lopinavir–ritonavir					such as D-dimer, CRP,	
						and IL-6	
21	Shufeng Jiedu capsule+WM vs.	RCS	34/34	Moderate	1) Improve the symptoms of cough, sputum, fatigue,	1) Increase lymphocyte	51
	WM				chest tightness, and shortness of breath	count	
					2) Lower the rate of transferring to severe disease	2) Decrease the levels of	

					3) Promote the absorption of lung inflammation and	CRP, procalcitonin, and	
					improve lung CT imaging	D-dimer	
22	Shufeng Jiedu capsule+arbidol vs.	RCS	100/100	Mild	1) Alleviate the symptoms of fever, cough, chest	Increase lymphocyte	52
	arbidol				distress, and shortness of breath	count and lymphocyte	
					2) Increase the absorption lung infected lesions	percentage	
23	Shufeng Jiedu capsule+arbidol vs.	RCS	40/30	Mild or	1) Shorten the antipyretic time and the disappearance	Unreported	53
	arbidol			moderate	time of dry cough, nasal congestion, runny nose,		
					pharyngeal pain, fatigue, and diarrhea		
					2) Reduce novel coronavirus negative conversion time		
24	Shufeng Jiedu capsule+arbidol vs.	RCS	100/100	Moderate	1) Shorten defervescence time	1) Increase WBC and	54
	arbidol				2) Improve resolution of pneumonia on chest CT	lymphocyte count	
						2) Reduce the levels of	
						CRP and IL-6	
25	Hanshiyi formula+WM vs. WM	RCS	430/291	Mild or	Reduce the progression to severe disease	Unreported	55
				moderate			
26	Lianhua Qingke granules+WM	RCT	25/32	Mild or	Ameliorate the symptoms of cough, sputum, fever,	Unreported	56
	vs. WM			moderate	fatigue, dry throat, and sore throat, and shorten the		
					duration of cough and sputum, reduce lung diseases,		
					improve respiratory function		
27	Toujie Quwen	RCS	32/33	Mild or	1) Improve the symptoms of fever, cough, fatigue,	1) Up-regulate	57
	granules+moxifloxacin+ambroxol			moderate	expectoration, dry throat, and sore throat	lymphocyte count and	
	vs. moxifloxacin+ambroxol				2) Improve lung CT imaging	neutrophil ratio	
						2) Down-regulate the	
						levels of CRP, D-dimer,	
						and procalcitonin	
28	Reyanning mixture+WM vs. WM	Multi-center	26/23	Moderate	1) Improve the symptoms of dry throat, cough, fatigue,	No significant	58
		clinical			chest tightness, and headache, and shorten the duration	differences in neutrophil	
		observation			of fever	count, lymphocyte count	
					2) Promote the improvement of lung CT	and CRP level	

					3) Improve nucleic acid negative conversion rate				
29	Maxing Shigan decoction+WM	Before and	40/0	Moderate	Improve the symptoms of fever, cough, fatigue,	Decrease CRP level,	59		
		after			hemoptysis, nausea, vomiting, diarrhea, and chest pain	increase CD4 ⁺ T and			
		comparison				CD8 ⁺ T count			
30	Honeysuckle oral liquid+WM vs.	Multi-center	200/100	Moderate	1) Shorten the length of hospitalization and the time of	No significant	60		
	WM	clinical			nucleic acid negative conversion	difference in the levels			
		observation			2) Lower right lung CT score	of ALT, AST,			
						creatinine, and uric acid			
31	Chansu injection+WM vs. WM	RCT	25/25	Severe or	Improve the respiratory function and shorten the	Improve the respiratory	61		
				critical	respiratory support step-down time	function indicators of			
						PaO ₂ /FiO ₂ and ROX			
						index			
32	Yidu-toxicity blocking lung	RCT	15/24	Severe	All patients are cured and discharged	Reduce the levels of	62		
	decoction+WM vs. WM					IL-6 and TNF-α			
33	Qingfei Dayuan granules+WM	Multi-center	451/0	All	1) Reduce the incidence of fever, cough, and fatigue	1) Increase lymphocyte	63		
		clinical			2) Improve the symptoms of aversion to cold, nasal	count			
		observation			obstruction, runny nose, sneezing, pharyngeal itch, sore	2) Reduce the levels of			
					throat, dyspnea, chest tightness, muscle ache or joint	CRP and procalcitonin			
					pain, dizziness, headache, tolerance, nausea and				
					vomiting, abdominal distension, and loose stool				
					3) Thin white greasy moss, thick greasy moss, and				
					yellow greasy moss, and improve tongue color				
					4) Decrease and thin lung lesion area				
34	"Fei Yan No. 1"+WM vs. WM	RCS	49/35	All	1) Improve the rate of recovering from symptoms and	Reduce leukocyte count	64		
					shorten the time	and CRP level			
					2) Increase the proportion of testing negative for nucleic				
					acid				
					3) Promote focal lung absorption and inflammation				
35	Xuanfei Huazhuo decoction+WM	Case analysis	40/0	All	1) Improve the symptoms of cough, fever, sputum,	1) Improve WBC count,	65		

					diarrhea, loss of appetite, and fatigue	lymphocyte, and	
					2) Promote the absorption of pulmonary inflammation	neutrophil percentage	
						2) Reduce the levels of	
						CRP, ESR, total	
						bilirubin, LDH, and the	
						ratio of AST/ALT	
36	Keguan-1+WM vs. WM	RCT	24/24	All	1) Reduce ARDS development	No significant	66
					2) Shorten the time to fever resolution	difference in	
					3) Tend to improve lung injury recovery	biochemical indicators	
						such as ALT, AST, and	
						D-dimer	
37	Qingfei Touxie Fuzheng	RCT	51/49	All	1) Alleviate the symptoms of fever, cough,	Decrease the levels of	67
	recipe+WM vs. WM				expectoration, chest tightness, and shortness of breath	ESR, CRP, and IL-6,	
					2) Promote the absorption of pulmonary lesions and	tend to increase IFN-γ	
					improve oxygenation	level	
38	Ganlu Xiaodu decoction+Chinese	Case analysis	131/0	All	Increase the effective rate of lung lesions absorption	Increase WBC and	68
	medicine and WM					lymphocyte count	
39	Matrine injection+WM	Case analysis	40/0	All	1) Improve the symptoms of cough, fatigue, appetite,	Alleviate absolute value	69
					and digestive tract	and ratio of lymphocyte	
					2) Promote absorption of lung lesions, especially for	and CRP	
					grid-like and fibrotic lesions		
					3) Shorten nucleic acid clearance time		
40	Diammonium	Case analysis	46/0	All	Improve the symptoms of low-grade fever, cough, and	1) Increase lymphocyte	70
	glycyrrhizinate+arbidol				fatigue	count and decrease ESR	
						level	
						2) Decrease the levels of	
						CRP, IL-6, and	
						procalcitonin	

Table 2 Clinical evidence of TCM for typical characteristics of COVID-19.

Clinical evidence	TCM
Clinical symptom	
Fever	Jinhua Qinggan granules ³⁷ , Lianhua Qingwen capsule ^{27,28,38–40} , Qingfei Paidu decoction ^{46,47} , Xuanfei Baidu Decoction ³² ,
	Shufeng Jiedu capsule ^{52,53} , Lianhua Qingke granules ⁵⁶ , Toujie Quwen granules ⁵⁷ , Reyanning mixture ⁵⁸ , Maxing Shigan
	decoction ⁵⁹ , Qingfei Dayuan granules ⁶³ , Xuanfei Huazhuo decoction ⁶⁵ , Qingfei Touxie Fuzheng recipe ⁶⁷ , Diammonium
	glycyrrhizinate ⁷⁰
Cough	Jinhua Qinggan granules ³⁷ , Lianhua Qingwen capsule ^{27,38–40} , Qingfei Paidu decoction ^{46,47} , Xuanfei Baidu Decoction ³² , Shufeng
	Jiedu capsule ^{51–53} , Lianhua Qingke granules ⁵⁶ , Toujie Quwen granules ⁵⁷ , Reyanning mixture ⁵⁸ , Maxing Shigan decoction ⁵⁹ ,
	Qingfei Dayuan granules ⁶³ , Xuanfei Huazhuo decoction ⁶⁵ , Qingfei Touxie Fuzheng recipe ⁶⁷ , Matrine injection ⁶⁹ ,
	Diammonium glycyrrhizinate ⁷⁰
Fatigue	Jinhua Qinggan granules ³⁷ , Lianhua Qingwen capsule ^{27,28,39,40} , Qingfei Paidu decoction ⁴⁶ , Xuanfei Baidu Decoction ³² , Shufeng
	Jiedu capsule ^{51,53} , Lianhua Qingke granules ⁵⁶ , Toujie Quwen granules ⁵⁷ , Reyanning mixture ⁵⁸ , Maxing Shigan decoction ⁵⁹ ,
	Qingfei Dayuan granules ⁶³ , Xuanfei Huazhuo decoction ⁶⁵ , Matrine injection ⁶⁹ , Diammonium glycyrrhizinate ⁷⁰
Dry throat	Lianhua Qingwen capsule ³⁹ , Shufeng Jiedu capsule ⁵³ , Lianhua Qingke granules ⁵⁶ , Toujie Quwen granules ⁵⁷ , Reyanning
	mixture ⁵⁸
Sore throat	Lianhua Qingwen capsule ³⁹ , Lianhua Qingke granules ⁵⁶ , Toujie Quwen granules ⁵⁷ , Qingfei Dayuan granules ⁶³
Sputum production	Jinhua Qinggan granules ³⁷ , Qingfei Paidu decoction ⁴⁹ , Lianhua Qingke granules ⁵⁶ , Xuanfei Huazhuo decoction ⁶⁵ , Qingfei
	Touxie Fuzheng recipe ⁶⁷
Shortness of breath	Lianhua Qingwen capsule ^{28,38} , Qingfei Paidu decoction ⁴⁶ , Shufeng Jiedu capsule ^{51,52} , Qingfei Dayuan granules ⁶³ , Qingfei
	Touxie Fuzheng recipe ⁶⁷
Myalgia	Lianhua Qingwen capsule ²⁸ , Shufeng Jiedu capsule ⁵³ , Qingfei Dayuan granules ⁶³
Diarrhea	Lianhua Qingwen capsule ²⁸ , Shufeng Jiedu capsule ⁵³ , Maxing Shigan decoction ⁵⁹ , Xuanfei Huazhuo decoction ⁶⁵
Duration of nucleic	Jinhua Qinggan granules ³⁶ , Lianhua Qingwen capsule ⁴¹ , Qingfei Paidu decoction ⁴⁸ , Shufeng Jiedu capsule ⁵³ , Reyanning
acid turn negative	mixture ⁵⁸ , Honeysuckle oral liquid ⁶⁰ , "Fei Yan No. 1" ⁶⁴ , Matrine injection ⁶⁹
Time to symptom	Lianhua Qingwen capsule ²⁷ , Xuebijing injection ⁴⁴ , Qingfei Paidu decoction ^{47–50} , Huashi Baidu Decoction ³¹ , Honeysuckle oral
recovery	liquid ⁶⁰ , Yidu-toxicity blocking lung decoction ⁶² , "Fei Yan No. 1" ⁶⁴ , Keguan-1 ⁶⁶
The progression to	Shufeng Jiedu capsule ⁵¹ , Hanshiyi formula ⁵⁵

severe disease

Multiorgan injury Xuebijing injection⁴⁴, Qingfei Paidu decoction⁴⁸

Lung feature

Lung inflammatory Jinhua Qinggan granules³⁶, Lianhua Qingwen capsule⁴¹, Xuebijing injection⁴⁴, Qingfei Paidu decoction⁴⁵, Shufeng Jiedu

absorption capsule⁵², "Fei Yan No. 1"⁶⁴, Xuanfei Huazhuo decoction⁶⁵, Ganlu Xiaodu decoction⁶⁸, Matrine injection⁶⁹

CT imaging Lianhua Qingwen capsule²⁷, Qingfei Paidu decoction^{46,47}, Huashi Baidu formula³⁰, Shufeng Jiedu capsule⁵¹, Toujie Quwen

granules⁵⁷, Reyanning mixture⁵⁸, Honeysuckle oral liquid⁶⁰

Lung injury Xuebijing injection⁴², Lianhua Qingke granules⁵⁶, Qingfei Dayuan granules⁶³, Keguan-1⁶⁶

Lung function Chansu injection⁶¹

Laboratory finding

WBC count Jinhua Qinggan granules³⁶, Lianhua Qingwen capsule³⁹, Xuebijing injection^{43,44}, Xuanfei Baidu Decoction³², Shufeng Jiedu

capsule54, "Fei Yan No. 1"64, Xuanfei Huazhuo decoction65, Ganlu Xiaodu decoction68

Lymphocyte count Jinhua Qinggan granules³⁶, Lianhua Qingwen capsule^{39,41}, Xuebijing injection^{43,44}, Qingfei Paidu decoction^{45,46}, Xuanfei Baidu

Decoction³², Shufeng Jiedu capsule⁵¹, Toujie Quwen granules⁵⁷, Qingfei Dayuan granules⁶³, Xuanfei Huazhuo decoction⁶⁵,

Ganlu Xiaodu decoction⁶⁸, Matrine injection⁶⁹, Diammonium glycyrrhizinate⁷⁰

Oxygenation index Xuebijing injection⁴³, Chansu injection⁶¹

CRP Lianhua Qingwen capsule³⁹, Xuebijing injection^{43,44}, Qingfei Paidu decoction⁴⁵, Xuanfei Baidu Decoction³², Huashi Baidu

formula³⁰, Shufeng Jiedu capsule^{51,54}, Toujie Quwen granules⁵⁷, Maxing Shigan decoction⁵⁹, Qingfei Dayuan granules⁶³, "Fei

Yan No. 1"64, Xuanfei Huazhuo decoction65, Qingfei Touxie Fuzheng recipe67, Matrine injection69, Diammonium

glycyrrhizinate70

IL-6 Xuebijing injection⁴², Qingfei Paidu decoction⁴⁸, Shufeng Jiedu capsule⁵⁴, Yidu–toxicity blocking lung decoction⁶², Qingfei

Touxie Fuzheng recipe⁶⁷, Diammonium glycyrrhizinate⁷⁰

TNF-α Xuebijing injection⁴², Yidu–toxicity blocking lung decoction⁶²

ESR Xuebijing injection⁴⁴, Qingfei Paidu decoction⁴⁶, Xuanfei Baidu Decoction³², Huashi Baidu formula³⁰, Xuanfei Huazhuo

decoction⁶⁵, Qingfei Touxie Fuzheng recipe⁶⁷, Diammonium glycyrrhizinate⁷⁰

CK Qingfei Paidu decoction⁴⁵

LDH Qingfei Paidu decoction⁴⁵, Xuanfei Huazhuo decoction⁶⁵

ALT Qingfei Paidu decoction⁴⁶, Xuanfei Huazhuo decoction⁶⁵

AST Qingfei Paidu decoction⁴⁶, Xuanfei Huazhuo decoction⁶⁵

Procalcitonin	Lianhua Qingwen capsule ³⁹ , Shufeng Jiedu capsule ⁵¹ , Toujie Quwen granules ⁵⁷ , Qingfei Dayuan granules ⁶³ , Diammonium
	glycyrrhizinate ⁷⁰
D-dimer	Qingfei Paidu decoction ⁴⁶ , Shufeng Jiedu capsule ⁵¹ , Toujie Quwen granules ⁵⁷
CD4 ⁺ T cell	Maxing Shigan decoction ⁵⁹
CD8 ⁺ T cell	Maxing Shigan decoction ⁵⁹

Table 3 Potential mechanisms of TCM for COVID-19.

No.	TCM	Coronavirus	Model/method	IC ₅₀ (EC ₅₀) or	Potential mechanism	Ref.
				dosage		
1	Jinhua Qinggan	SARS-CoV-2	Network	Not applicable	1) Regulate TNF, PI3K/Akt, and HIF-1 signaling	88
			pharmacology (NP),	(NA)	pathways via binding angiotensin converting	
			molecular docking		enzyme 2 (ACE2) and acting on targets such as	
					PTGS2, HSP90AB1, HSP90AA1, PTGS1, and	
					NCOA2	
					2) Formononetin, stigmasterol, β -sitosterol, and	
					anhydroicaritin have a high affinity with 3CLpro	
					and ACE2	
2	Lianhua Qingwen	SARS-CoV-2	Infected Vero E6	411.2 μg/mL	1) Inhibit virus replication and decrease the number	89
	capsule		cells and Huh-7		of virus particles	
			cells, cytopathic		2) Reduce pro-inflammatory cytokines of TNF- α ,	
			effect (CPE), plaque		IL-6, MCP-1, and IP-10 production	
			reduction assay			
3	Lianhua Qingwen	SARS-CoV-2	NP	NA	1) Exert antiviral effect and repair lung injury	90
	formula				2) Modulate inflammatory process and relieve	
					cytokine storm	
					3) Improve ACE2 expression disorder caused	
					symptoms	
4	Xuebijing injection	SARS-CoV-2	Infected Vero E6	11.75 mg/mL	1) Exert antiviral effect and reduce plaque	42
			cells and Huh-7		formation	

			cells, CPE, plaque		2) Inhibit the expression and release of TNF- α ,	
			reduction assay		IL-6, MIP-1 β , RANTES, and IP-10	
5	Xuebijing injection	SARS-CoV-2	NP, molecular	NA	1) Quercetin, luteolin, apigenin, and other	91
			docking		compounds may target TNF, MAPK1, and IL6	
					2) Anhydrosafflor yellow B, salvianolic acid B, and	
					rutin play the role of anti-inflammatory, antiviral,	
					and immune response	
6	Xuebijing injection	SARS-CoV-2	NP	NA	Exert anti-inflammatory and immunoregulatory	92
					effects through RAS, NF-κB, PI3K, Akt, MAPK,	
					VEGF, TLR, TNF, and TRP signaling pathways	
7	Qingfei Paidu	SARS-CoV-2	NP, molecular	NA	1) Exert antiviral and anti-inflammatory activities,	93–95
	decoction		docking		regulate metabolic programming, and repair lung	
					injury	
					2) Glycyrrhizin in one of the main ingredients	
					inhibits TLR agonists induced IL-6 production in	
					macrophage	
8	Qingfei Paidu	SARS-CoV-2	NP, molecular	NA	1) Exhibit the effects of immune regulation,	96
	decoction		docking, molecular		anti-infection, anti-inflammation, and multi-organ	
			verification		protection	
					2) Four compounds of baicalin, glycyrrhizin,	
					hesperidin, and hyperoside act on the targets	
					including AKT1, TNF-α, IL-6, PTGS2, HMOX1,	
					IL10, and TP53	
					3) Inhibit IL-6, CCL2, TNF- α , NF- κ B, PTGS1/2,	
					CYP1A1, and CYP3A4 activity, and increase IL-10	
					expression	
					4) Reduce platelet aggregation.	
9	Huashi Baidu	SARS-CoV-2	NP, molecular	NA	1) Regulate TNF, PI3K-Akt, NOD-like, MAPK,	97
	formula		docking		and HIF-1 signaling pathways	

					2) Baicalein and quercetin are the top two	
					compounds with a high affinity to ACE2	
10	Xuanfei Baidu	SARS-CoV-2	NP	NA	Regulate viral, parasites and bacterial infections,	98
					and modulate energy metabolism, immunity, and	
					inflammation	
11	Shufeng Jiedu	SARS-CoV-2	NP	NA	Regulate the key targets of RELA, MAPK1,	99
	capsule				MAPK14, CASP3, CASP8, and IL-6	
12	Shufeng Jiedu	SARS-CoV-2	NP, molecular	NA	Regulate immunomodulatory and	100
	capsule		docking		anti-inflammatory related targets on multiple	
					pathways	
13	Maxing Shigan	SARS-CoV-2	NP	NA	1) Reduce inflammation and suppress cytokine	101
	decoction				storm	
					2) Protect pulmonary alveolar-capillary barrier and	
					alleviate pulmonary edema	
					3) Regulate immune response and decrease fever	
14	Maxing Shigan	SARS-CoV-2	NP, molecular	NA	1) Inhibit IL-6 mediated JAK-STAT signal	102
	decoction		docking, molecular		pathway	
			verification		2) Amygdalin is predicted to bind ACE2, 3CLpro,	
					and RdRp	
15	Cold-damp plague	SARS-CoV-2	NP, molecular	NA	1) Regulate free radical production and blood	103
	formula		docking		circulation	
					2) Exert antiviral, immune- regulatory, and	
					anti- inflammatory by targeting ACE2 and IL-6	
16	Dayuanyin	SARS-CoV-2	NP, molecular	NA	1) Play an anti-inflammatory and	104,105
			docking		immunoregulatory role via acting on IL-6, IL-1 β ,	
					and CCL2	
					2) Decrease the level of IL6 in mild, moderate, and	
					severe clinical cases	
					3) The ingredients of kaempferol, quercetin,	

					7-methoxy-2-methyl, isoflavone, naringenin, and	
					formononetin target IL-6, IL-1 β , and CCL2 with	
					high affinity	
17	Reduning injection	SARS-CoV-2	Infected Vero E6	103.420 μg/mL	Exert antiviral effect	106
	2 3		cells, CPE, NP	1.5	2) Regulate ACE2, 3CLpro, and PLpro activity	
			, ,		3) Modulate inflammation-related expressions of	
					MAPKs, PKC, and NF-κB	
18	Liu Shen capsule	SARS-CoV-2	Infected Vero E6	0.6 μg/mL	Inhibit virus replication and reduce plaque	107
			cells and Huh-7	****	formation	
			cells, CPE, plaque		 Reduce pro-inflammatory cytokines of TNF-α, 	
			reduction assay		IL-6, IL-1 β , IL-8, MCP-1, and IP-10 production,	
			reduction assay		and inhibit p-NF- κ B p65, p-I κ B α , and p-p38	
					MAPK expression	
19	Pudilan Xiaoyan	SARS-CoV-2	Infected Vero E6	1.08 mg/mL		108
19	•	SARS-COV-2		1.06 mg/mL	Inhibit viral replication in vitro and in vivo	108
20	oral liquid		cells, CPE	1) 0.02 1.2		100
20	Shuanghuanglian	SARS-CoV-2	1) Infected Vero	1) 0.93–1.2	1) Inhibit viral replication	109
	preparation		cells	μL/mL	2) Inhibit 3CLpro activity	
			2) Enzyme inhibition	2) 0.06–0.09		
			assay	μL/mL		
21	Yinqiao powder	SARS-CoV-2	NP, molecular	NA	Regulate TNF, T-cell receptor, Toll-like receptor,	110
			docking, surface		and MAPK signaling pathways	
			plasmon resonance			
			(SPR) analysis			
22	Pudilan prescription	SARS-CoV-2	NP, GSEA	NA	1) Prevent SARS-CoV-2 entrance by blocking	111
			enrichment,		ACE2	
			molecular docking		2) Inhibit cytokine storm of CRP, IFN-γ, IL-6,	
					IL-10, TNF, EGFR, CCL5, and TGF- β 1	
23	Matrine injection	SARS-CoV-2	NP, molecular	NA	1) Inhibit viral replication, host cell apoptosis and	112,113
			docking		inflammation by targeting the TNF- α , IL-6, and	

					CASP3 in TNF signaling pathway	
					2) Reduce lung tissue damage and lung index	
					3) Decrease the production of IL-6, IL-10, TNF- α ,	
					IFN- γ , as well as the viral load in lung tissue	
					4) Increase the percentage of CD4 ⁺ T cells, CD8 ⁺ T	
					cells and B cells in peripheral blood	
24	Shenfu decoction	SARS-CoV-2	NP, molecular	NA	Play antiviral role through multi-component,	114
			docking		multi-target, and multi-pathway approach, and	
					exert anti-inflammation, immune regulation, and	
					multi-organ protection effects	
25	Andrographis	SARS-CoV-2	Infected Calu-3	$0.036~\mu g/mL$	Exert antiviral effect	115
	paniculate		cells, CPE			
	(Chuanxinlian)					
26	Scutellaria	SARS-CoV-2	1) Enzyme inhibition	1) 8.52 mg/mL	1) Inhibit 3CLpro activity	116
	baicalensis		assay	2) 0.74 mg/mL	2) Exert antiviral effect	
	(Huangqin)		2) Infected Vero			
			cells, CPE			
27	Rheum officinale	SARS-CoV	Infected Vero E6	$1{-}10~\mu g/mL$	Block spike-ACE2 interaction	117
	(Yaoyong		cells, CPE,			
	Dahuang)		biotinylated ELISA			
28	Polygonum	SARS-CoV	Infected Vero E6	$1-10~\mu g/mL$	Block spike–ACE2 interaction	117
	multiflorum		cells, CPE,			
	(Heshouwu)		Biotinylated ELISA			
29	Houttuynia cordata	SARS-CoV	Flow cytometry,	$0800~\mu\text{g/mL}$	1) Stimulate the proliferation of mouse splenic	118
	(Yuxingcao)		ELISA, enzyme		lymphocytes	
			inhibition assay, etc.		2) Increase the proportion of CD4 $^{\scriptscriptstyle +}$ and CD8 $^{\scriptscriptstyle +}$ T	
					cells	
					3) Increase in the secretion of IL-2 and IL-10 in	
					mouse splenic lymphocytes	

					4) Inhibit 3CLpro and RdRp activity	
30	Rheum palmatum	SARS-CoV	Enzyme inhibition	13.76 μg/mL	Inhibit 3CLpro activity	119
	(Zhangye Dahuang)		assay			
31	Cibotium barometz	SARS-CoV	1) Infected Vero E6	1) 8.42 μg/mL	1) Inhibit viral replication	120
	(Gouji)		cells, CPE	2) 39 μg/mL	2) Inhibit 3CLpro activity	
			2) Enzyme inhibition			
			assay			
32	Dioscorea batatas	SARS-CoV	1) Infected Vero E6	1) 8.06 μg/mL	1) Inhibit viral replication	120
	(Shanyao)		cells, CPE	2) 44 μg/mL	2) Inhibit 3CLpro activity	
			2) Enzyme inhibition			
			assay			

Table 4 Potential mechanisms of TCM ingredients for COVID-19.

No.	TCM ingredient	Source	Coronavirus	Model/method	$IC_{50}(EC_{50})$	Potential mechanism	Ref.
					or dosage		
1	Rhein	Rheum palmatum	SARS-CoV-2	Enzyme inhibition	18.33	Inhibit ACE2 activity	121
		(Yaoyong Dahuang)		assay, molecular	μmol/L		
				docking, and surface			
				plasmon resonance			
				(SPR) analysis			
2	Forsythoside A	Forsythiae fructus	SARS-CoV-2	Enzyme inhibition	Unclear	Inhibit ACE2 activity	121
		(Lianqiao) fruit		assay, molecular			
				docking, SPR			
				analysis			
3	Neochlorogenic acid	Lonicera japonica	SARS-CoV-2	Enzyme inhibition	\sim 40 μ mol/L	Inhibit ACE2 activity	121
		(Jingyinhua)		assay, molecular			
				docking, SPR			
				analysis			
4	Quercetin	Ginkgo biloba	SARS-CoV-2	Enzyme inhibition	4.48 μmol/L	Inhibit ACE2 activity	122

		(Yingxing)		assay			
5	Ephedrine	Ephedrae Herba	SARS-CoV-2	Molecular docking,	Unclear	Inhibit ACE2 activity	123
		(Mahuang)		SPR analysis			
6	Hesperidin	Citrus aurantium	SARS-CoV-2	Target-based virtual	Unclear	Block spike-ACE2	124,125
		(Suancheng)		ligand screening		interaction.	
7	Geniposide	Gardenia jasminoides	SARS-CoV-2	Molecular docking	Unclear	Inhibit TMPRSS2 activity	126
		(Zhizi)					
8	Baicalin	Scutellaria baicalensis	SARS-CoV-2	1) Infected Vero E6	1) 27.87	1) Inhibit viral replication	109
		(Huangqin)		cells, CPE	μmol/L	2) Inhibit 3CLpro activity	
				2) Enzyme	2) 6.41		
				inhibition assay	μmol/L		
9	Baicalein	Scutellaria baicalensis	SARS-CoV-2	1) Enzyme	1) 0.39	1) Inhibit 3CLpro activity	116
		(Huangqin)		inhibition assay	μmol/L	2) Exert antiviral infection	
				2) Infected Vero	2) 2.9	effect	
				cells	μmol/L		
10	Shikonin	Lithospermum	SARS-CoV-2	Enzyme inhibition	15.75	Inhibit 3CLpro activity	127
		erythrorhizon (Zicao)		assay	μmol/L		
11	EGCG	Green tea	SARS-CoV-2	Enzyme inhibition	0.017	Inhibit 3CLpro activity	128
				assay	μmol/L		
12	Theaflavin	Black tea	SARS-CoV-2	Enzyme inhibition	0.015	Inhibit 3CLpro activity	128
				assay	μmol/L		
13	Scutellarein	Scutellaria baicalensis	SARS-CoV-2	Enzyme inhibition	5.8 μmol/L	Inhibit 3CLpro activity	116
		(Huangqin)		assay			
14	Myricetin	Myrica rubra (Yangmei)	SARS-CoV-2	Enzyme inhibition	2.86 μmol/L	Inhibit 3CLpro activity	116
				assay			
15	Cannabidiol	Cannabis sativa (Dama)	SARS-CoV-2	1) Molecular	7.91 μmol/L	1) Bind to PLpro	129
				docking		2) Exert antiviral effect	
				2) Infected Vero			
				cells			

16	Theaflavin	Black tea	SARS-CoV-2	Molecular docking	Unclear	Inhibit RdRp activity	130,131
17	Digitoxin	Digitalis purpurea	SARS-CoV-2	Infected Vero cells,	$0.23~\mu mol/L$	Exert antiviral effect	132
		(Yangdihuang)		СРЕ			
18	Tetrandrine	Stephania tetrandra	SARS-CoV-2	Infected Vero cells,	3 μmol/L	Exert antiviral effect	132
		(Fengfangji)		СРЕ			
19	Glycyrrhizin	Glycyrrhiza uralensis	SARS-CoV-2	Infected Vero E6	$0.53~\mu mol/L$	Exert antiviral effect	133
		(Gancao)		cells, CPE			
20	Resveratrol	Polygonum cuspidatum	SARS-CoV-2	Infected Vero E6,	66 μmol/L	Exert antiviral effect	134
		(Huzhang)		Calu-3 and primary			
				human bronchial			
				epithelium cells,			
				СРЕ			
21	Pterostilbene	Pterocarpus santalinus	SARS-CoV-2	Infected Vero E6,	19 μmol/L	Exert antiviral effect	134
		(Zitan)		Calu-3 and primary			
				human bronchial			
				epithelium cells,			
				СРЕ			
22	Phillyrin	Forsythiae fructus	SARS-CoV-2	Infected Vero-E6	1) 63.9	1) Inhibit viral replication	135
		(Lianqiao)		cells and Huh-7	$\mu g/mL$	2) Reduce the production	
				cells, CPE	2) and 3)	of proinflammatory	
					62.5–250	cytokines of TNF-α, IL-6,	
					μg/mL	IL-1 β , MCP-1, and IP-10	
						3) Suppress NF-κB	
						signaling pathway	
23	Catechin	Green tea	SARS-CoV-2	Molecular docking	Unclear	Bind to 3CLpro, cathepsin	131,136
						L, RBD of S protein,	
						NSP6, and nucleocapsid	
						protein	
24	Artemisinin	Artemisia annua	SARS-CoV-2	Infected Vero E6	64.45	Inhibit viral replication	137

		(Qinghao)		cells, CPE	μmol/L		
25	Artesunate	Artemisinin derivative	SARS-CoV-2	Infected Vero E6	12.98	Inhibit viral replication	137
				cells, CPE	μmol/L		
26	Cepharanthine	Stephania japonica	SARS-CoV-2	Infected Vero E6	0.98 μmol/L	Inhibit viral entry and viral	138
		(Qianjinteng)		cells, CPE		replication	
27	Bufalin	Toad venom (Chansu)	SARS-CoV-2	Infected Vero E6	18 nmol/L	Exert antiviral effect by	139
				cells, CPE		targeting Na ⁺ /K ⁺ -ATPase	
28	Bruceine A	Brucea javanica	SARS-CoV-2	Infected Vero E6	11 nmol/L	Exert antiviral effect	139
		(Yadanzi)		cells, CPE			
29	Naringenin	Gardenia jasminoides	SARS-CoV-2	Infected Vevo E6	31.3–250	Target two-pore channel 2	140
		(Zhishi)		cells, CPE	μmol/L		
30	Andrographolide	Andrographis paniculate	SARS-CoV-2	Infected Calu-3	0.034	Exert antiviral effect	115
		(Chuanxinlian)		cells, CPE	μmol/L		
31	Glycyrrhizin+vitamin	Glycyrrhiza uralensis	SARS-CoV-2	NP	Unclear	Elevate immunity and	141
	С	(Gancao)				suppress inflammatory	
						stress	
32	Chlorogenic acid	Lonicera japonica	SARS-CoV-2	NP	Unclear	Exert antiviral effect by	142
		(Jinyinhua)				targeting NFE2L2,	
						PPARG, ESR1, ACE, IL6,	
						and HMOX1	
33	Emodin	Rheum palmatum	SARS-CoV	Infected Vero E6	200 μmol/L	Block spike–ACE2	117
		(Yaoyong Dahuang)		cells, CPE,		interaction	
				biotinylated ELISA			
34	Celastrol	Celastrus orbiculatus	SARS-CoV	Enzyme inhibition	10.3 μmol/L	Inhibit 3CLpro activity	143,144
		(Nansheteng)		assay			
35	Tingenone	Euonymus alatus	SARS-CoV	Enzyme inhibition	9.9 μmol/L	Inhibit 3CLpro activity	143
		(Weimao)		assay			
36	Curcurmin	Curcuma longa	SARS-CoV	1) Enzyme	1) 23.5	1) Inhibit 3CLpro activity	145,146
		(Jianghuang)		inhibition assay;	μmol/L	2) Inhibit viral replication	

				2) Infected Vero E6	2) 40		
				cells, CPE	μmol/L		
37	Quercetin	Ginkgo biloba	SARS-CoV	Enzyme inhibition	73 μmol/L	Inhibit 3CLpro activity	147,148
		(Yingxing)		assay			
38	Tanshinone IIA	Salvia miltiorrhiza	SARS-CoV	Enzyme inhibition	89.1 μmol/L	Inhibit 3CLpro activity	149
		(Danshen)		assay			
39	Dihydrotanshinone I	Salvia miltiorrhiza	SARS-CoV	Enzyme inhibition	14.4 μmol/L	Inhibit 3CLpro activity	149
		(Danshen)		assay			
40	Xanthoangelol E	Angelica keiskei	SARS-CoV	Enzyme inhibition	11.4 μmol/L	Inhibit 3CLpro activity	150
		(Mingriye)		assay			
41	Sinigrin	Isatis indigotica root	SARS-CoV	Enzyme inhibition	217 μmol/L	Inhibit 3CLpro activity	151
		(Banlangen)		assay			
42	Hesperetin	Isatis indigotica root	SARS-CoV	Enzyme inhibition	8.3 μmol/L	Inhibit 3CLpro activity	151
		(Banlangen)		assay			
43	Pectolinarin	Cirsium japonicum	SARS-CoV	Enzyme inhibition	37.78	Inhibit 3CLpro activity	152
		(Daji)		assay	μmol/L		
44	Luteolin	(Jinyinhua)	SARS-CoV	1) Infected Vero E6	1) 9.02	1) Exert antiviral effect	153,154
				cells, CPE;	μmol/L	2) Inhibit 3CLpro activity	
				2) Enzyme	2) 20.2		
				inhibition assay	μmol/L		
45	Hirsutenone	Alnus japonica	SARS-CoV	Enzyme inhibition	4.1 μmol/L	Inhibit PLpro activity	155
		(Chiyang)		assay			
46	Tanshinone IIB	Salvia miltiorrhiza	SARS-CoV	Enzyme inhibition	10.7 μmol/L	Inhibit PLpro activity	149
		(Danshen)		assay			
47	Crytotanshinone	Salvia miltiorrhiza	SARS-CoV	Enzyme inhibition	0.8 μmol/L	Inhibit PLpro activity	149
		(Danshen)		assay			
48	Dihydrotanshinone I	Salvia miltiorrhiza	SARS-CoV	Enzyme inhibition	4.9 μmol/L	Inhibit PLpro activity	149
		(Danshen)		assay			
49	Xanthoangelol E	Angelica keiskei	SARS-CoV	Enzyme inhibition	1.2 μmol/L	Inhibit PLpro activity	150

		(Mingriye)		assay			
50	Terrestrimine	Tribulus terrestris	SARS-CoV	Enzyme inhibition	15.8 μmol/L	Inhibit PLpro activity	156
		(Cijili) fruits		assay			
51	Isobavachalcone	Psoralea corylifolia	SARS-CoV	Enzyme inhibition	7.3 µmol/L	Inhibit PLpro activity	157
		(Buguzhi) seeds		assay			
52	Psoralidin	Psoralea corylifolia	SARS-CoV	Enzyme inhibition	$4.2~\mu mol/L$	Inhibit PLpro activity	157
		(Buguzhi) seeds		assay			
53	Tomentin A-E	Paulownia tomentosa	SARS-CoV	Enzyme inhibition	5.0–12.5	Inhibit PLpro activity	158
		fruits (Maopaotong)		assay	μmol/L		
54	Glycyrrhizin	Glycyrrhiza uralensis	SARS-CoV	Infected Vero cells,	0.3 mg/mL	Inhibit virus replication	159–161
		(Gancao)		CPE			
55	Cepharanthine	Stephania japonica	SARS-CoV	Infected Vero E6	6.0-9.5	Exert antiviral effect	162
		(Qianjinteng)		cells, CPE	$\mu g/mL$		
56	Ginsenoside Rb1	Panax ginseng	SARS-CoV	Infected Vero E6	$100 \ \mu mol/L$	Exert antiviral effect	163
		(Renshen)		cells, CPE			
57	Aescin	Aesculus chinensis	SARS-CoV	Infected Vero E6	6.0 μmol/L	Inhibit viral replication	163
		(Qiyeshu)		cells, CPE			
58	Reserpine	Ophiorrhiza japonica	SARS-CoV	Infected Vero E6	$3.4 \ \mu mol/L$	Inhibit viral replication	163
		(Shegencao)		cells, CPE			
59	Lycorine	Lycoris radiata	SARS-CoV	Infected Vero E6	15.7 nmol/L	Exert antiviral effect	164
		(Shisuan)		cells, CPE			